

TENT COOPERATION TRE Y

PCT
NOTIFICATION OF ELECTION
(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

Date of mailing (day/month/year) 14 June 2001 (14.06.01)	To: Commissioner US Department of Commerce United States Patent and Trademark Office, PCT 2011 South Clark Place Room CP2/5C24 Arlington, VA 22202 ETATS-UNIS D'AMERIQUE in its capacity as elected Office
International application No. PCT/US00/24722	Applicant's or agent's file reference 2026-4316PC
International filing date (day/month/year) 08 September 2000 (08.09.00)	Priority date (day/month/year) 10 September 1999 (10.09.99)
Applicant CUTTITTA, Frank et al	
<p>1. The designated Office is hereby notified of its election made:</p> <p><input checked="" type="checkbox"/> in the demand filed with the International Preliminary Examining Authority on: 03 April 2001 (03.04.01)</p> <p><input type="checkbox"/> in a notice effecting later election filed with the International Bureau on: _____</p> <p>2. The election <input checked="" type="checkbox"/> was <input type="checkbox"/> was not</p> <p>made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).</p>	

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland Facsimile No.: (41-22) 740.14.35	Authorized officer Claudio Borton Telephone No.: (41-22) 338.83.38
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PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference 2026-4316PC	FOR FURTHER ACTION	see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.
International application No. PCT/US 00/24722	International filing date (day/month/year) 08/09/2000	(Earliest) Priority Date (day/month/year) 10/09/1999
Applicant		

THE GOVERNMENT OF THE UNITED STATES OF AMERICA as

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 4 sheets.

It is also accompanied by a copy of each prior art document cited in this report.

1. Basis of the report

- a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.
 - the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).
- b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :
 - contained in the international application in written form.
 - filed together with the international application in computer readable form.
 - furnished subsequently to this Authority in written form.
 - furnished subsequently to this Authority in computer readable form.
 - the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
 - the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. **Certain claims were found unsearchable** (See Box I).

3. **Unity of invention is lacking** (see Box II).

4. With regard to the **title**,

- the text is approved as submitted by the applicant.
- the text has been established by this Authority to read as follows:

DETERMINATION OF ADRENOMEDULLIN-BINDING PROTEINS

5. With regard to the **abstract**,

- the text is approved as submitted by the applicant.
- the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No.

- as suggested by the applicant.
- because the applicant failed to suggest a figure.
- because this figure better characterizes the invention.

1

None of the figures.

INTERNATIONAL SEARCH REPORT

International Application No

US 00/24722

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 G01N33/74 C07K16/00 A61K38/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, BIOSIS, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MARTINEZ ALFREDO ET AL: "Regulation of insulin secretion and blood glucose metabolism by adrenomedullin." ENDOCRINOLOGY, vol. 137, no. 6, 1996, pages 2626-2632, XP000982198 ISSN: 0013-7227 the whole document ---	1-52
A	WO 97 07214 A (UNSWORTH EDWARD J ;MACRI CHARLES (US); US HEALTH (US); CUTTITTA FR) 27 February 1997 (1997-02-27) the whole document --- -/-	1-52

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

16 February 2001

Date of mailing of the international search report

26/02/2001

Name and mailing address of the ISA

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Authorized officer

Pellegrini, P

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/24722

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MARTINEZ ALFREDO ET AL: "Expression of adrenomedullin and its receptor in normal and malignant human skin: A potential pluripotent role in the integument." ENDOCRINOLOGY, vol. 138, no. 12, December 1997 (1997-12), pages 5597-5604, XP000982199 ISSN: 0013-7227 cited in the application the whole document ---	1-52
A	US 4 883 784 A (KANEKO ISAO) 28 November 1989 (1989-11-28) the whole document ---	51, 52
P, A	MARTINEZ ALFREDO ET AL: "Adrenomedullin binding protein-1 (AMBP-1) modulates tumor growth by reducing availability of adrenomedullin (AM)." PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page 153 XP000982163 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000 ISSN: 0197-016X abstract ---	1-52
P, A	PIO RUBEN ET AL: "Isolation and characterization of a human plasma adrenomedullin binding protein. Analysis of its expression in human tumor cell lines." PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page 152 XP000982162 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000 ISSN: 0197-016X abstract ---	1-52

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/24722

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,A	<p>ELSASSER TED H ET AL: "Adrenomedullin binding protein in the plasma of multiple species: Characterization by radioligand blotting." ENDOCRINOLOGY, vol. 140, no. 10, October 1999 (1999-10), pages 4908-4911, XP000982200 ISSN: 0013-7227 cited in the application the whole document</p> <p>---</p>	1-52
P,A	<p>MARTINEZ A ET AL: "Is adrenomedullin a causal agent in some cases of type 2 diabetes?" PEPTIDES (NEW YORK)., vol. 20, no. 12, December 1999 (1999-12), pages 1471-1478, XP000982202 ISSN: 0196-9781 the whole document</p> <p>-----</p>	1-52

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

US 00/24722

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 9707214	A 27-02-1997	AT 180832	T	15-06-1999
		AU 710662	B	23-09-1999
		AU 6776596	A	12-03-1997
		CA 2229741	A	27-02-1997
		DE 69602756	D	08-07-1999
		DE 69602756	T	10-02-2000
		EP 0845036	A	03-06-1998
		JP 11512087	T	19-10-1999
US 4883784	A 28-11-1989	AT 61227	T	15-03-1991
		CA 1294872	A	28-01-1992
		DE 3677897	D	11-04-1991
		EP 0222611	A	20-05-1987
		JP 2004739	C	20-12-1995
		JP 7042235	B	10-05-1995
		JP 62201822	A	05-09-1987

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

by fax and post

To:

FEILER, William S.
Morgan & Finnegan, L.L.P.
345 Park Avenue
New York, New York 10154
ETATS-UNIS D'AMERIQUE

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2001 DEC 21 AID 51

MORGAN & FINNAGAN LLP

NOTIFICATION OF TRANSMITTAL OF
THE INTERNATIONAL PRELIMINARY
EXAMINATION REPORT

(PCT Rule 71.1)

Date of mailing
(day/month/year)

14.12.2001

Applicant's or agent's file reference

2026-4316PC ✓

IMPORTANT NOTIFICATION

International application No/
PCT/US00/24722 ✓

International filing date (day/month/year)
08/09/2000 ✓

Priority date (day/month/year)
10/09/1999 ✓

Applicant

THE GOVERNMENT OF THE UNITED STATES OF AMERICA as

1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.
2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

4. REMINDER

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/

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Authorized officer

MYLONAS, E

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PATENT COOPERATION TREATY
PCT
INTERNATIONAL PRELIMINARY EXAMINATION REPORT
(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 2026-4316PC	FOR FURTHER ACTION		See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
International application No. PCT/US00/24722	International filing date (day/month/year) 08/09/2000	Priority date (day/month/year) 10/09/1999	
International Patent Classification (IPC) or national classification and IPC G01N33/68			
<p>Applicant THE GOVERNMENT OF THE UNITED STATES OF AMERICA as</p>			
<p>1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of 10 sheets, including this cover sheet.</p> <p><input type="checkbox"/> This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).</p> <p>These annexes consist of a total of sheets.</p>			
<p>3. This report contains indications relating to the following items:</p> <ul style="list-style-type: none"> I <input checked="" type="checkbox"/> Basis of the report II <input checked="" type="checkbox"/> Priority III <input checked="" type="checkbox"/> Non-establishment of opinion with regard to novelty, inventive step and industrial applicability IV <input type="checkbox"/> Lack of unity of invention V <input checked="" type="checkbox"/> Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement VI <input type="checkbox"/> Certain documents cited VII <input checked="" type="checkbox"/> Certain defects in the international application VIII <input checked="" type="checkbox"/> Certain observations on the international application 			

Date of submission of the demand 03/04/2001	Date of completion of this report 14.12.2001
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer  Vanmontfort, D Telephone No. +49 89 2399 8457

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 2026-4316PC	FOR FURTHER ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)	
International application No. PCT/US00/24722	International filing date (day/month/year) 08/09/2000	Priority date (day/month/year) 10/09/1999
International Patent Classification (IPC) or national classification and IPC G01N33/68		
<p>Applicant</p> <p>THE GOVERNMENT OF THE UNITED STATES OF AMERICA as</p> <p>1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of 10 sheets, including this cover sheet.</p> <p><input type="checkbox"/> This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).</p> <p>These annexes consist of a total of sheets.</p> <p>3. This report contains indications relating to the following items:</p> <ul style="list-style-type: none"> I <input checked="" type="checkbox"/> Basis of the report II <input checked="" type="checkbox"/> Priority III <input checked="" type="checkbox"/> Non-establishment of opinion with regard to novelty, inventive step and industrial applicability IV <input type="checkbox"/> Lack of unity of invention V <input checked="" type="checkbox"/> Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement VI <input type="checkbox"/> Certain documents cited VII <input checked="" type="checkbox"/> Certain defects in the international application VIII <input checked="" type="checkbox"/> Certain observations on the international application 		

Date of submission of the demand 03/04/2001	Date of completion of this report 14.12.2001
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Vanmontfort, D Telephone No. +49 89 2399 8457



**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US00/24722

I. Basis of the report

1. With regard to the **elements** of the international application (*Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)*):
Description, pages:

1-58 as originally filed

Claims, No.:

1-52 as originally filed

Drawings, sheets:

1/18-18/18 as originally filed

Sequence listing part of the description, pages:

1-2, as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- the language of publication of the international application (under Rule 48.3(b)).
- the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- contained in the international application in written form.
- filed together with the international application in computer readable form.
- furnished subsequently to this Authority in written form.
- furnished subsequently to this Authority in computer readable form.
- The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US00/24722

the description, pages:

the claims, Nos.:

the drawings, sheets:

5. This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):
(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:

II. Priority

1. This report has been established as if no priority had been claimed due to the failure to furnish within the prescribed time limit the requested:

copy of the earlier application whose priority has been claimed.

translation of the earlier application whose priority has been claimed.

2. This report has been established as if no priority had been claimed due to the fact that the priority claim has been found invalid.

Thus for the purposes of this report, the international filing date indicated above is considered to be the relevant date.

3. Additional observations, if necessary:
see separate sheet

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:

the entire international application.

claims Nos. 24, 25, 41-52.

because:

the said international application, or the said claims Nos. 24, 25, 41-52 with respect to industrial applicability relate to the following subject matter which does not require an international preliminary examination (specify):
see separate sheet

the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US00/24722

that no meaningful opinion could be formed (*specify*):

- the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.
- no international search report has been established for the said claims Nos. .

2. A meaningful international preliminary examination cannot be carried out due to the failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions:

- the written form has not been furnished or does not comply with the standard.
- the computer readable form has not been furnished or does not comply with the standard.

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes: Claims 1-23,26-50, 52
	No: Claims 24, 25, 51
Inventive step (IS)	Yes: Claims 1-22, 26-50
	No: Claims 23-25, 51, 52

Industrial applicability (IA) Yes: Claims 1-23, 26-40
No: Claims

2. Citations and explanations
see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:
see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:
see separate sheet

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/24722

1. Section II

Claims 1-5, 15, 16, 21-25, 37-40, 45-47, 51 and 52 of the present application are not entitled to the claimed priority date (10.09.1999). There is no disclosure in the priority document of an improved quantitative assay for detecting AM in serum utilizing a chaotropic agent (claims 1-5, 15 and 16), a method of identifying antagonist agents that inhibit AM, factor H or AM/fH activity (claims 21 and 22), the use of said antagonists in a method of treating cancer or a condition associated with decreased secretion of insulin (claims 23-25, 45-47), a kit comprising an antibody against the AM/fH complex (claims 37-40) or the use of factor H in a method of treating a condition associated with an autoimmune response to pancreatic cells (claims 51 and 52).

2. Section III

Claims 24, 25 and 41-52 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(i) PCT).

3. Section V

Reference is made to the following documents:

- D1 MARTINEZ et al., ENDOCRINOLOGY, vol. 138, no. 12, December 1997 (1997-12), pages 5597-5604, XP000982199
- D2 US-A-4 883 784 (KANEKO ISAO) 28 November 1989
- D3 MARTINEZ et al., ENDOCRINOLOGY, vol. 137, no. 6, 1996, pages 2626-2632, XP000982198
- D4 MARTINEZ et al., PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000, page 153 XP000982163 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000
- D5 PIO RUBEN et al., PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/24722

152 XP000982162 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000

D6 ELSASSER et al., ENDOCRINOLOGY, vol. 140, no. 10, October 1999 (1999-10), pages 4908-4911, XP000982200, cited in the application
D7 MARTINEZ et al., PEPTIDES (NEW YORK)., vol. 20, no. 12, December 1999, pages 1471-1478, XP000982202

D4-D7 are indicated in the search report as P-documents. Since claims 1-5, 15, 16, 21-25, 37-40, 45-47, 51 and 52 of the present application are not entitled to the claimed priority date (10.09.1999), D4-D7 could be relevant for these claims with respect to novelty and inventive step (Articles 33(2) and (3) PCT).

3.1 The subject-matter of claims 1-5 is novel and inventive (Articles 33(2) and 33(3) PCT).
D1 (page 5598, right column, first paragraph), which is considered to represent the most relevant state of the art, discloses a RIA to measure AM levels in a sample. The subject-matter of claim 1 differs from D1 in that prior to serum extraction and quantification of AM, the sample is treated with a chaotropic agent to dissociate AM from factor H. The problem to be solved by the present invention may therefore be regarded as the provision of a more accurate method to measure AM. There is no indication in any of the available prior art documents that the routine RIA quantification of AM, as described in D1, fails to account for the amount of AM bound to its binding protein, factor H. Furthermore, the proposed solution (pretreating the sample with a chaotropic agent to dissociate AM from factor H) is also not disclosed. Therefore, it would not be obvious for a person skilled in the art to introduce this feature in the closest prior art to solve the problem posed. Consequently, the subject-matter of claim 1 is considered to involve an inventive step. The same applies to dependent claims 2-5.

3.2 The subject-matter of claims 6-16 is novel and inventive (Articles 33(2) and 33(3) PCT).
None of the available prior art documents discloses or refers to a complex comprising AM and the newly identified AM binding protein, human complement factor H. Hence, the use of factor H as an AM-binding protein in a method to

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

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monitor the AM concentrations in a subject is novel and inventive. Therefore, the subject-matter of claim 6 is novel and inventive. The same applies to dependent claims 7-16.

3.3 The subject-matter of claims 17-20 is novel and inventive (Articles 33(2) and 33(3) PCT).

There is no disclosure or reference in any of the available prior art documents of a kit comprising factor H and at least one component to facilitate or detect AM-binding. Therefore, the subject-matter of claim 17 is novel and inventive. The same applies to dependent claims 18-20.

3.4 The subject-matter of claims 21 and 22 is novel and inventive (Articles 33(2) and 33(3) PCT).

Claims 21 and 22 are directed against a method of identifying an antagonist agent which inhibits the complexation of adrenomedullin and factor H. There is no indication in any of the available prior art documents of a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, it would not be obvious for one skilled in the art to identify antagonists which inhibits the formation of the above-mentioned complex. Hence, the subject-matter of claims 21 and 22 is novel and inventive.

3.5 The subject-matter of claim 23 is not inventive (Article 33(3) PCT).

For the assessment of claims of products defined in terms of a process of identification, a product is not rendered novel merely by the fact that it is identified by means of a new process how difficult or time-consuming the technology may be. Claims for products defined in terms of a process of identification are admissible only if the products as such fulfil the requirements of patentability, i.e. that they are novel and inventive.

D4-D6 disclose the existence of an AM-binding protein, such as AMBP-1. Said binding protein would have been identified by the method of claim 21.

Consequently, as it is common practice to provide an antagonist agent in the form of a composition, the subject-matter of claim 23 can not be considered to involve an inventive step (Article 33(3) PCT).

3.6 The subject-matter of claims 24 and 25 is not novel (Articles 33(2) PCT).

**INTERNATIONAL PRELIMINARY
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International application No. PCT/US00/24722

D4 and D5 disclose the existence of an AM-binding protein, such as AMBP-1, which is known to modulate tumor growth. Since said binding protein would have been identified by the method of claim 21, D4 and D5 anticipate the novelty of the subject-matter of claims 24 and 25.

3.7 The subject-matter of claim 26 is novel and inventive (Articles 33(2) and 33(3) PCT).

None of the available prior art documents discloses or refers to a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, the subject-matter of claim 26 is novel and inventive.

3.8 The subject-matter of claims 27 and 28 is novel and inventive (Articles 33(2) and 33(3) PCT).

D3 (page 2626, right column, paragraph antibodies), which is considered to represent the most relevant state of the art, discloses a monoclonal antibody MoAb-G6 which is directed against the PO72 peptide of AM. The problem to be solved by the present invention may therefore be regarded as the provision of an alternative antibody against AM. There is no indication in any of the available prior art documents of a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, it would not be obvious for one skilled in the art to design antibodies against said novel complex. Hence, the subject-matter of claims 27 and 28 is novel and inventive.

3.9 The subject-matter of claims 29-44 and 48-50 is novel and inventive (Articles 33(2) and 33(3) PCT).

The use of the antibody of claim 27 in a pharmaceutical composition (claim 29), in a method of isolating AM/fH complex from a biological sample (claims 30 and 31), in a method of detecting AM/fH complex in a biological sample (claims 32-36), in a kit (claims 37-40), in a method of inhibiting the growth or proliferation of cancer or tumor cells (claims 41-44) and in a method of treating a condition associated with decreased secretion of insulin by pancreatic cells (claims 48-50) is novel and inventive, given that the antibody of claim 27 is novel and inventive.

3.10 The subject-matter of claims 45-47 is novel and inventive (Articles 33(2) and 33(3) PCT).

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/24722

There is no disclosure or reference in any of the available prior art documents that a substance which inhibits the complexation of AM and factor H can be used for the treatment of a condition associated with decreased secretion of insulin by pancreatic cells. Therefore, the subject-matter of claims 45-47 is novel and inventive.

3.11 The subject-matter of claim 51 is not novel (Articles 33(2) PCT). D4 discloses the use of factor H in the therapy of autoimmune diseases. Therefore, D4 anticipates the novelty of the subject-matter of claim 51.

3.12 Dependent claim 52 does not contain any features which, in combination with the features of any claim to which it refers, meet the requirements of the PCT in respect of inventive step (Article 33(3) PCT). The feature is merely one of several straightforward possibilities from which the skilled person would select, in accordance with circumstances, without the exercise of inventive skill.

3.13 For the assessment of the present claims 24, 25 and 41-52 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.

4. Section VII

4.1 Page 1 lines 6-9 is irrelevant subject-matter according to Rule 9.1 (iv) PCT.

4.2 The vague and imprecise statement "without departing from the scope and spirit of the invention" in the description (page 58 lines 9-13) implies that the subject-matter for which protection is sought may be different to that defined by the claims, thereby resulting in lack of clarity (Article 6 PCT) when used to interpret them (see also the PCT Guidelines, C-III, 4.3a).

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/24722

4.3 Contrary to the requirements of Rule 5.1(a)(ii) PCT, the relevant background art disclosed in D2, D4 and D5 is not mentioned in the description, nor are these documents identified therein.

5. Section VIII

Claims 24, 25 and 41-52 are directed to methods of treating cancer or diabetes. The application does not meet the requirements of Article 6 PCT because none of these claims enjoy any substantial support in terms of concrete examples in the application as filed. There is no evidence whatsoever in the application as filed that the suggested methods actually yield the desired result.

PATENT COOPERATION TREATY

2026-4316 PC
Dehoratius

From the:
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To:

FEILER, William S.
Morgan & Finnegan, L.L.P.
345 Park Avenue
New York, New York 10154
ETATS-UNIS D'AMERIQUE

PTO/PCT Rec'd 08 MAR 2002

PCT

WRITTEN OPINION

(PCT Rule 66)

Date of mailing (day/month/year)	08.08.2001
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Applicant's or agent's file reference 2026-4316PC	REPLY DUE	within 3 month(s) from the above date of mailing
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International application No. PCT/US00/24722	International filing date (day/month/year) 08/09/2000	Priority date (day/month/year) 10/09/1999
---	--	--

International Patent Classification (IPC) or both national classification and IPC G01N33/68
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Applicant

THE GOVERNMENT OF THE UNITED STATES OF AMERICA as

1. This written opinion is the first drawn up by this International Preliminary Examining Authority.

2. This opinion contains indications relating to the following items:

- I Basis of the opinion
- II Priority
- III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV Lack of unity of invention
- V Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI Certain document cited
- VII Certain defects in the international application
- VIII Certain observations on the international application

3. The applicant is hereby invited to reply to this opinion.

When? See the time limit indicated above. The applicant may, before the expiration of that time limit, request this Authority to grant an extension, see Rule 66.2(d).

How? By submitting a written reply, accompanied, where appropriate, by amendments, according to Rule 66.3. For the form and the language of the amendments, see Rules 66.8 and 66.9.

Also: For an additional opportunity to submit amendments, see Rule 66.4. For the examiner's obligation to consider amendments and/or arguments, see Rule 66.4 bis. For an informal communication with the examiner, see Rule 66.6.

If no reply is filed, the international preliminary examination report will be established on the basis of this opinion.

4. The final date by which the international preliminary examination report must be established according to Rule 69.2 is: 10/01/2002.

CASE 2026-4316 PC ATTY COH
DUE November 8, 2001 (wr. opn)
1 mo. call-up October 8, 2001
BY J. M.

Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 eprmu d Fax: +49 89 2399 - 4465

Authorized officer / Examiner

Vanmontfort, D

Formalities officer (incl. extension of time limits) Danti, B Telephone No. +49 89 2399 8161
--



I. Basis of the opinion

1. With regard to the **elements** of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this opinion as "originally filed"):

Description, pages:

1-58 as originally filed

Claims, No.:

1-52 as originally filed

Drawings, sheets:

1/18-18/18 as originally filed

Sequence listing part of the description, pages:

1-2, as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- the language of publication of the international application (under Rule 48.3(b)).
- the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- contained in the international application in written form.
- filed together with the international application in computer readable form.
- furnished subsequently to this Authority in written form.
- furnished subsequently to this Authority in computer readable form.
- The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

the description, pages:
 the claims, Nos.:
 the drawings, sheets:

5. This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:**II. Priority**

1. This opinion has been established as if no priority had been claimed due to the failure to furnish within the prescribed time limit the requested:
 - copy of the earlier application whose priority has been claimed.
 - translation of the earlier application whose priority has been claimed.
2. This opinion has been established as if no priority had been claimed due to the fact that the priority claim has been found invalid.

Thus for the purposes of this opinion, the international filing date indicated above is considered to be the relevant date.

3. Additional observations, if necessary:
see separate sheet

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been and will not be examined in respect of:
 - the entire international application,
 - claims Nos. 24, 25, 41-52,

because:

- the said international application, or the said claims Nos. 24, 25, 41-52 with respect to industrial applicability relate to the following subject matter which does not require an international preliminary examination (specify):
see separate sheet

WRITTEN OPINION

International application No. PCT/US00/24722

- the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear that no meaningful opinion could be formed (*specify*):
- the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.
- no international search report has been established for the said claims Nos. .

2. A written opinion cannot be drawn due to the failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions:

- the written form has not been furnished or does not comply with the standard.
- the computer readable form has not been furnished or does not comply with the standard.

V. Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Claims 24, 25, 51
Inventive step (IS)	Claims 23-25, 51, 52
Industrial applicability (IA)	Claims

2. Citations and explanations
see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:
see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:
see separate sheet

1. Section II

Claims 1-5, 15, 16, 21-25, 37-40, 45-47, 51 and 52 of the present application are not entitled to the claimed priority date (10.09.1999). There is no disclosure in the priority document of an improved quantitative assay for detecting AM in serum utilizing a chaotropic agent (claims 1-5, 15 and 16), a method of identifying antagonist agents that inhibit AM, factor H or AM/fH activity (claims 21 and 22), the use of said antagonists in a method of treating cancer or a condition associated with decreased secretion of insulin (claims 23-25, 45-47), a kit comprising an antibody against the AM/fH complex (claims 37-40) or the use of factor H in a method of treating a condition associated with an autoimmune response to pancreatic cells (claims 51 and 52).

2. Section III

Claims 24, 25 and 41-52 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(i) PCT).

3. Section V

Reference is made to the following documents:

- D1 MARTINEZ et al., ENDOCRINOLOGY, vol. 138, no. 12, December 1997 (1997-12), pages 5597-5604, XP000982199
- D2 US-A-4 883 784 (KANEKO ISAO) 28 November 1989
- D3 MARTINEZ et al., ENDOCRINOLOGY, vol. 137, no. 6, 1996, pages 2626-2632, XP000982198
- D4 MARTINEZ et al., PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000, page 153 XP000982163 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000
- D5 PIO RUBEN et al., PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page

152 XP000982162 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000

D6 ELSASSER et al., ENDOCRINOLOGY, vol. 140, no. 10, October 1999 (1999-10), pages 4908-4911, XP000982200, cited in the application
D7 MARTINEZ et al., PEPTIDES (NEW YORK), vol. 20, no. 12, December 1999, pages 1471-1478, XP000982202

D4-D7 are indicated in the search report as P-documents. Since claims 1-5, 15, 16, 21-25, 37-40, 45-47, 51 and 52 of the present application are not entitled to the claimed priority date (10.09.1999), D4-D7 could be relevant for these claims with respect to novelty and inventive step (Articles 33(2) and (3) PCT).

3.1 The subject-matter of claims 1-5 is novel and inventive (Articles 33(2) and 33(3) PCT).

D1 (page 5598, right column, first paragraph), which is considered to represent the most relevant state of the art, discloses a RIA to measure AM levels in a sample. The subject-matter of claim 1 differs from D1 in that prior to serum extraction and quantification of AM, the sample is treated with a chaotropic agent to dissociate AM from factor H. The problem to be solved by the present invention may therefore be regarded as the provision of a more accurate method to measure AM. There is no indication in any of the available prior art documents that the routine RIA quantification of AM, as described in D1, fails to account for the amount of AM bound to its binding protein, factor H. Furthermore, the proposed solution (pretreating the sample with a chaotropic agent to dissociate AM from factor H) is also not disclosed. Therefore, it would not be obvious for a person skilled in the art to introduce this feature in the closest prior art to solve the problem posed. Consequently, the subject-matter of claim 1 is considered to involve an inventive step. The same applies to dependent claims 2-5.

3.2 The subject-matter of claims 6-17 is novel and inventive (Articles 33(2) and 33(3) PCT).

None of the available prior art documents discloses or refers to a complex comprising AM and the newly identified AM binding protein, human complement factor H. Hence, the use of factor H as an AM-binding protein in a method to

monitor the AM concentrations in a subject is novel and inventive. Therefore, the subject-matter of claim 6 is novel and inventive. The same applies to dependent claims 7-16.

3.3 The subject-matter of claims 17-20 is novel and inventive (Articles 33(2) and 33(3) PCT).

There is no disclosure or reference in any of the available prior art documents of a kit comprising factor H and at least one component to facilitate or detect AM-binding. Therefore, the subject-matter of claim 17 is novel and inventive. The same applies to dependent claims 18-20.

3.4 The subject-matter of claims 21 and 22 is novel and inventive (Articles 33(2) and 33(3) PCT).

Claims 21 and 22 are directed against a method of identifying an antagonist agent which inhibits the complexation of adrenomedullin and factor H. There is no indication in any of the available prior art documents of a complex comprising AM and the newly identified AM binding protein, human complement factor H.

Therefore, it would not be obvious for one skilled in the art to identify antagonists which inhibits the formation of the above-mentioned complex. Hence, the subject-matter of claims 21 and 22 is novel and inventive.

3.5 The subject-matter of claim 23 is not inventive (Article 33(3) PCT).

For the assessment of claims of products defined in terms of a process of identification, a product is not rendered novel merely by the fact that it is identified by means of a new process how difficult or time-consuming the technology may be. Claims for products defined in terms of a process of identification are admissible only if the products as such fulfil the requirements of patentability, i.e. that they are novel and inventive.

D4-D6 disclose the existence of an AM-binding protein, such as AMBP-1. Said binding protein would have been identified by the method of claim 21.

Consequently, as it is common practice to provide an antagonist agent in the form of a composition, the subject-matter of claim 23 can not be considered to involve an inventive step (Article 33(3) PCT).

3.6 The subject-matter of claims 24 and 25 is not novel (Articles 33(2) PCT).

D4 and D5 disclose the existence of an AM-binding protein, such as AMBP-1, which is known to modulate tumor growth. Since said binding protein would have been identified by the method of claim 21, D4 and D5 anticipate the novelty of the subject-matter of claims 24 and 25.

3.7 The subject-matter of claim 26 is novel and inventive (Articles 33(2) and 33(3) PCT).

None of the available prior art documents discloses or refers to a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, the subject-matter of claim 26 is novel and inventive.

3.8 The subject-matter of claims 27 and 28 is novel and inventive (Articles 33(2) and 33(3) PCT).

D3 (page 2626, right column, paragraph antibodies), which is considered to represent the most relevant state of the art, discloses a monoclonal antibody MoAb-G6 which is directed against the PO72 peptide of AM. The problem to be solved by the present invention may therefore be regarded as the provision of an alternative antibody against AM. There is no indication in any of the available prior art documents of a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, it would not be obvious for one skilled in the art to design antibodies against said novel complex. Hence, the subject-matter of claims 27 and 28 is novel and inventive.

3.9 The subject-matter of claims 29-44 and 48-50 is novel and inventive (Articles 33(2) and 33(3) PCT).

The use of the antibody of claim 27 in a pharmaceutical composition (claim 29), in a method of isolating AM/fH complex from a biological sample (claims 30 and 31), in a method of detecting AM/fH complex in a biological sample (claims 32-36), in a kit (claims 37-40), in a method of inhibiting the growth or proliferation of cancer or tumor cells (claims 41-44) and in a method of treating a condition associated with decreased secretion of insulin by pancreatic cells (claims 48-50) is novel and inventive, given that the antibody of claim 27 is novel and inventive.

3.10 The subject-matter of claims 45-47 is novel and inventive (Articles 33(2) and 33(3) PCT).

There is no disclosure or reference in any of the available prior art documents that a substance which inhibits the complexation of AM and factor H can be used for the treatment of a condition associated with decreased secretion of insulin by pancreatic cells. Therefore, the subject-matter of claims 45-47 is novel and inventive.

3.11 The subject-matter of claim 51 is not novel (Articles 33(2) PCT). D4 discloses the use of factor H in the therapy of autoimmune diseases. Therefore, D4 anticipates the novelty of the subject-matter of claim 51.

3.12 Dependent claim 52 does not contain any features which, in combination with the features of any claim to which it refers, meet the requirements of the PCT in respect of inventive step (Article 33(3) PCT). The feature is merely one of several straightforward possibilities from which the skilled person would select, in accordance with circumstances, without the exercise of inventive skill.

3.13 For the assessment of the present claims 24, 25 and 41-52 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.

4. Section VII

4.1 Page 1 lines 6-9 is irrelevant subject-matter according to Rule 9.1 (iv) PCT.

4.2 The vague and imprecise statement "without departing from the scope and spirit of the invention" in the description (page 58 lines 9-13) implies that the subject-matter for which protection is sought may be different to that defined by the claims, thereby resulting in lack of clarity (Article 6 PCT) when used to interpret them (see also the PCT Guidelines, C-III, 4.3a).

4.3 Contrary to the requirements of Rule 5.1(a)(ii) PCT, the relevant background art disclosed in D2, D4 and D5 is not mentioned in the description, nor are these documents identified therein.

5. Section VIII

Claims 24, 25 and 41-52 are directed to methods of treating cancer or diabetes. The application does not meet the requirements of Article 6 PCT because none of these claims enjoy any substantial support in terms of concrete examples in the application as filed. There is no evidence whatsoever in the application as filed that the suggested methods actually yield the desired result.

PATENT COOPERATION TREATY

2026-4316 PC

C. Dehorter

PCT
COMMUNICATION

NOTICE INFORMING THE APPLICANT OF THE COMMUNICATION OF THE INTERNATIONAL APPLICATION TO THE DESIGNATED OFFICES

(PCT Rule 47.1(c), first sentence)
PTO/PCT Rec'd

To:

FEILER, William, S.
Morgan & Finnegan, L.L.P.
345 Park Avenue
New York, NY 10154
ETATS-UNIS D'AMERIQUE

08 MAR 2002

Date of mailing (day/month/year)
15 March 2001 (15.03.01)

Applicant's or agent's file reference
2026-4316PC

IMPORTANT NOTICE

International application No.	International filing date (day/month/year)
PCT/US00/24722	08 September 2000 (08.09.00)

Priority date (day/month/year)

10 September 1999 (10.09.99)

Applicant	THE GOVERNMENT OF THE UNITED STATES OF AMERICA as represented by THE SECRETARY, DEPARTMENT OF HEALTH et al
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1. Notice is hereby given that the International Bureau has communicated, as provided in Article 20, the international application to the following designated Offices on the date indicated above as the date of mailing of this Notice:

AU,KP,KR,US

In accordance with Rule 47.1(c), third sentence, those Offices will accept the present Notice as conclusive evidence that the communication of the international application has duly taken place on the date of mailing indicated above and no copy of the international application is required to be furnished by the applicant to the designated Office(s).

2. The following designated Offices have waived the requirement for such a communication at this time:

AE,AG,AL,AM,AP,AT,AZ,BA,BB,BG,BR,BY,BZ,CA,CH,CN,CR,CU,CZ,DE,DK,DM,DZ,EA,EE,EP,ES,
FI,GB,GD,GE,GH,GM,HR,HU,IL,IN,IS,JP,KE,KG,KZ,LC,LK,LR,LS,LT,LU,LV,MA,MD,MG,MK,
MN,MW,MX,MZ,NO,NZ,OA,PL,PT,RO,RU,SD,SE,SG,SI,SK,SL,TJ,TM,TR,TT,TZ,UA,UG,UZ,VN,YU,
The communication will be made to those Offices only upon their request. Furthermore, those Offices do not require the applicant to furnish a copy of the international application (Rule 49.1(a-bis)).

3. Enclosed with this Notice is a copy of the international application as published by the International Bureau on
15 March 2001 (15.03.01) under No. WO 01/18550

REMINDER REGARDING CHAPTER II (Article 31(2)(a) and Rule 54.2)

If the applicant wishes to postpone entry into the national phase until 30 months (or later in some Offices) from the priority date, a demand for international preliminary examination must be filed with the competent International Preliminary Examining Authority before the expiration of 19 months from the priority date.

It is the applicant's sole responsibility to monitor the 19-month time limit.

Note that only an applicant who is a national or resident of a PCT Contracting State which is bound by Chapter II has the right to file a demand for international preliminary examination.

REMINDER REGARDING ENTRY INTO THE NATIONAL PHASE (Article 22 or 39(1))

If the applicant wishes to proceed with the international application in the national phase, he must, within 20 months or 30 months, or later in some Offices, perform the acts referred to therein before each designated or elected Office.

For further important information on the time limits and acts to be performed for entering the national phase, see the Annex to Form PCT/IB/301 (Notification of Receipt of Record Copy) and Volume II of the PCT Applicant's Guide.

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland	Authorized officer J. Zahra
Facsimile No. (41-22) 740.14.35	Telephone No. (41-22) 338.83.38

PATENT COOPERATION TREATY

2026-4316 PC
Dehoratius.From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

PCT

To:

FEILER, William S.
 Morgan & Finnegan, L.L.P.
 345 Park Avenue
 New York, New York 10154
 ETATS-UNIS D'AMERIQUE

NOTIFICATION OF RECEIPT
OF DEMAND BY COMPETENT INTERNATIONAL
PRELIMINARY EXAMINING AUTHORITY(PCT Rules 59.3(e) and 61.1(b), first sentence
and Administrative Instructions, Section 601(a))Date of mailing
(day/month/year)

23.04.01

Applicant's or agent's file reference
2026-4316PC

IMPORTANT NOTIFICATION

International application No.
PCT/US 00/ 24722International filing date (day/month/year)
08/09/2000Priority date (day/month/year)
10/09/1999

Applicant

THE GOVERNMENT OF THE UNITED STATES OF AMERICA as

1. The applicant is hereby notified that this International Preliminary Examining Authority considers the following date as the date of receipt of the demand for international preliminary examination of the international application:

03/04/2001

2. This date of receipt is:

the actual date of receipt of the demand by this Authority (Rule 61.1(b)).
 the actual date of receipt of the demand on behalf of this Authority (Rule 59.3(e)).
 the date on which this Authority has, in response to the invitation to correct defects in the demand (Form PCT/IPEA/404), received the required corrections.

3. ATTENTION: That date of receipt is AFTER the expiration of 19 months from the priority date. Consequently, the election(s) made in the demand does (do) not have the effect of postponing the entry into the national phase until 30 months from the priority date (or later in some Offices) (Article 39(1)). Therefore, the acts for entry into the national phase must be performed within 20 months from the priority date (or later in some Offices) (Article 22). For details, see the *PCT Applicant's Guide*, Volume II.

(If applicable) This notification confirms the information given by telephone, facsimile transmission or in person on:

4. Only where paragraph 3 applies, a copy of this notification has been sent to the International Bureau.

Name and mailing address of the IPEA/



European Patent Office
 D-80298 Munich
 Tel. (+49-89) 2399-0, Tx: 523656 epmu d
 Fax: (+49-89) 2399-4465

Authorized officer

CHURCHER G

Tel. (+49-89) 2399-2545



PATENT COOPERATION TREATY

2026-4316 PC

From the INTERNATIONAL BUREAU

Dehoratios,

PCT

**NOTIFICATION CONCERNING
SUBMISSION OR TRANSMITTAL
OF PRIORITY DOCUMENT**

(PCT Administrative Instructions, Section 411)

Date of mailing (day/month/year)

14 December 2000 (14.12.00)

To:

2000 DEC 28 A 8:44

FEILER, William, S.
 Morgan & Finnegan LLP
 345 Park Avenue
 New York, NY 10154
 ETATS-UNIS D'AMERIQUE

Applicant's or agent's file reference

2026-4316PC ✓

IMPORTANT NOTIFICATION

International application No.

PCT/US00/24722 ✓

International filing date (day/month/year)

08 September 2000 (08.09.00) ✓

International publication date (day/month/year)

Not yet published

Priority date (day/month/year)

10 September 1999 (10.09.99) ✓

Applicant

THE GOVERNMENT OF THE UNITED STATES OF AMERICA as represented by THE
 SECRETARY, DEPARTMENT OF HEALTH et al

1. The applicant is hereby notified of the date of receipt (except where the letters "NR" appear in the right-hand column) by the International Bureau of the priority document(s) relating to the earlier application(s) indicated below. Unless otherwise indicated by an asterisk appearing next to a date of receipt, or by the letters "NR", in the right-hand column, the priority document concerned was submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b).
2. This updates and replaces any previously issued notification concerning submission or transmittal of priority documents.
3. An asterisk(*) appearing next to a date of receipt, in the right-hand column, denotes a priority document submitted or transmitted to the International Bureau but not in compliance with Rule 17.1(a) or (b). In such a case, the attention of the applicant is directed to Rule 17.1(c) which provides that no designated Office may disregard the priority claim concerned before giving the applicant an opportunity, upon entry into the national phase, to furnish the priority document within a time limit which is reasonable under the circumstances.
4. The letters "NR" appearing in the right-hand column denote a priority document which was not received by the International Bureau or which the applicant did not request the receiving Office to prepare and transmit to the International Bureau, as provided by Rule 17.1(a) or (b), respectively. In such a case, the attention of the applicant is directed to Rule 17.1(c) which provides that no designated Office may disregard the priority claim concerned before giving the applicant an opportunity, upon entry into the national phase, to furnish the priority document within a time limit which is reasonable under the circumstances.

Priority datePriority application No.Country or regional Office
or PCT receiving OfficeDate of receipt
of priority document

10 Sept 1999 (10.09.99)

60/153,397

US

11 Dec 2000 (11.12.00)

The International Bureau of WIPO
 34, chemin des Colibettes
 1211 Genève 20, Switzerland

Facsimile No. (41-22) 740.14.35

Authorized officer

Magda BOUACHA

3

Telephone No. (41-22) 338.83.38

PATENT COOPERATION TREATY

2026-436 PC

C. Dehoratius

From the INTERNATIONAL BUREAU

PCT

PTO/PCT Rec'd 08 MAR 2002

INFORMATION CONCERNING ELECTED OFFICES NOTIFIED OF THEIR ELECTION

(PCT Rule 61.3)

Date of mailing (day/month/year)
14 June 2001 (14.06.01)

To:
FEILER, William, S.
Morgan & Finnegan, L.L.P.
345 Park Avenue
New York, NY 10154
ETATS-UNIS D'AMERIQUE

Applicant's or agent's file reference
2026-4316PC

IMPORTANT INFORMATION

International application No. PCT/US00/24722	International filing date (day/month/year) 08 September 2000 (08.09.00)	Priority date (day/month/year) 10 September 1999 (10.09.99)
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Applicant THE GOVERNMENT OF THE UNITED STATES OF AMERICA as represented by THE SECRETARY, DEPARTMENT OF HEALTH et al

1. The applicant is hereby informed that the International Bureau has, according to Article 31(7), notified each of the following Offices of its election:

EP :AT,BE,CH,CY,DE,DK,ES,FI,FR,GB,GR,IE,IT,LU,MC,NL,PT,SE
National :AU,BG,CA,CN,CZ,DE,IL,JP,KP,KR,MN,NO,NZ,PL,RO,RU,SE,SK,US

2. The following Offices have waived the requirement for the notification of their election; the notification will be sent to them by the International Bureau only upon their request:

AP :GH,GM,KE,LS,MW,MZ,SD,SL,SZ,TZ,UG,ZW
EA :AM,AZ,BY,KG,KZ,MD,RU,TJ,TM
OA :BF,BJ,CF,CG,CI,CM,GA,GN,GW,ML,MR,NE,SN,TD,TG
National :AE,AG,AL,AM,AT,AZ,BA,BB,BR,BY,BZ,CH,CR,CU,DK,DM,DZ,EE,ES,FI,GB,
GD,GE,GH,GM,HR,HU,ID,IN,IS,KE,KG,KZ,LC,LK,LR,LS,LT,LU,LV,MA,MD,MG,MK,MW,
MX,MZ,PT,SD,SG,SI,SL,TJ,TM,TR,TT,TZ,UA,UG,UZ,VN,YU,ZA,ZW

3. The applicant is reminded that he must enter the "national phase" before the expiration of 30 months from the priority date before each of the Offices listed above. This must be done by paying the national fee(s) and furnishing, if prescribed, a translation of the international application (Article 39(1)(a)), as well as, where applicable, by furnishing a translation of any annexes of the international preliminary examination report (Article 36(3)(b) and Rule 74.1).

Some offices have fixed time limits expiring later than the above-mentioned time limit. For detailed information about the applicable time limits and the acts to be performed upon entry into the national phase before a particular Office, see Volume II of the PCT Applicant's Guide.

The entry into the European regional phase is postponed until 31 months from the priority date for all States designated for the purposes of obtaining a European patent.

The International Bureau of WIPO
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PATENT COOPERATION TREATY

PCT

REC'D 18 DEC 2001
WIPO PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70) 14

Applicant's or agent's file reference 2026-4316PC	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
International application No. PCT/US00/24722	International filing date (day/month/year) 08/09/2000	Priority date (day/month/year) 10/09/1999

International Patent Classification (IPC) or national classification and IPC
G01N33/68

Applicant

THE GOVERNMENT OF THE UNITED STATES OF AMERICA as

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 10 sheets, including this cover sheet.

This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

These annexes consist of a total of sheets.

3. This report contains indications relating to the following items:

- I Basis of the report
- II Priority
- III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV Lack of unity of invention
- V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI Certain documents cited
- VII Certain defects in the international application
- VIII Certain observations on the international application

Date of submission of the demand 03/04/2001	Date of completion of this report 14.12.2001
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Vanmontfort, D Telephone No. +49 89 2399 8457



**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US00/24722

I. Basis of the report

1. With regard to the **elements** of the international application (*Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)*):
Description, pages:

1-58 as originally filed

Claims, No.:

1-52 as originally filed

Drawings, sheets:

1/18-18/18 as originally filed

Sequence listing part of the description, pages:

1-2, as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- the language of publication of the international application (under Rule 48.3(b)).
- the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- contained in the international application in written form.
- filed together with the international application in computer readable form.
- furnished subsequently to this Authority in written form.
- furnished subsequently to this Authority in computer readable form.
- The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US00/24722

the description, pages:

the claims, Nos.:

the drawings, sheets:

5. This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)): *(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)*

6. Additional observations, if necessary:

II. Priority

1. This report has been established as if no priority had been claimed due to the failure to furnish within the prescribed time limit the requested:

copy of the earlier application whose priority has been claimed.

translation of the earlier application whose priority has been claimed.

2. This report has been established as if no priority had been claimed due to the fact that the priority claim has been found invalid.

Thus for the purposes of this report, the international filing date indicated above is considered to be the relevant date.

3. Additional observations, if necessary:
see separate sheet

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:

the entire international application.

claims Nos. 24, 25, 41-52.

because:

the said international application, or the said claims Nos. 24, 25, 41-52 with respect to industrial applicability relate to the following subject matter which does not require an international preliminary examination (specify):
see separate sheet

the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

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that no meaningful opinion could be formed (*specify*):

- the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.
- no international search report has been established for the said claims Nos. .

2. A meaningful international preliminary examination cannot be carried out due to the failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions:

- the written form has not been furnished or does not comply with the standard.
- the computer readable form has not been furnished or does not comply with the standard.

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes: Claims 1-23,26-50, 52
	No: Claims 24, 25, 51
Inventive step (IS)	Yes: Claims 1-22, 26-50
	No: Claims 23-25, 51, 52

Industrial applicability (IA) Yes: Claims 1-23, 26-40
No: Claims

2. Citations and explanations
see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:
see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:
see separate sheet

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/24722

1. Section II

Claims 1-5, 15, 16, 21-25, 37-40, 45-47, 51 and 52 of the present application are not entitled to the claimed priority date (10.09.1999). There is no disclosure in the priority document of an improved quantitative assay for detecting AM in serum utilizing a chaotropic agent (claims 1-5, 15 and 16), a method of identifying antagonist agents that inhibit AM, factor H or AM/fH activity (claims 21 and 22), the use of said antagonists in a method of treating cancer or a condition associated with decreased secretion of insulin (claims 23-25, 45-47), a kit comprising an antibody against the AM/fH complex (claims 37-40) or the use of factor H in a method of treating a condition associated with an autoimmune response to pancreatic cells (claims 51 and 52).

2. Section III

Claims 24, 25 and 41-52 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(i) PCT).

3. Section V

Reference is made to the following documents:

- D1 MARTINEZ et al., ENDOCRINOLOGY, vol. 138, no. 12, December 1997 (1997-12), pages 5597-5604, XP000982199
- D2 US-A-4 883 784 (KANEKO ISAO) 28 November 1989
- D3 MARTINEZ et al., ENDOCRINOLOGY, vol. 137, no. 6, 1996, pages 2626-2632, XP000982198
- D4 MARTINEZ et al., PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000, page 153 XP000982163 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000
- D5 PIO RUBEN et al., PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page

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EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/24722

152 XP000982162 91st Annual Meeting of the American Association for Cancer Research.;San Francisco, California, USA; April 01-05, 2000, March, 2000

D6 ELSASSER et al., ENDOCRINOLOGY, vol. 140, no. 10, October 1999 (1999-10), pages 4908-4911, XP000982200, cited in the application
D7 MARTINEZ et al., PEPTIDES (NEW YORK)., vol. 20, no. 12, December 1999, pages 1471-1478, XP000982202

D4-D7 are indicated in the search report as P-documents. Since claims 1-5, 15, 16, 21-25, 37-40, 45-47, 51 and 52 of the present application are not entitled to the claimed priority date (10.09.1999), D4-D7 could be relevant for these claims with respect to novelty and inventive step (Articles 33(2) and (3) PCT).

3.1 The subject-matter of claims 1-5 is novel and inventive (Articles 33(2) and 33(3) PCT).
D1 (page 5598, right column, first paragraph), which is considered to represent the most relevant state of the art, discloses a RIA to measure AM levels in a sample. The subject-matter of claim 1 differs from D1 in that prior to serum extraction and quantification of AM, the sample is treated with a chaotropic agent to dissociate AM from factor H. The problem to be solved by the present invention may therefore be regarded as the provision of a more accurate method to measure AM. There is no indication in any of the available prior art documents that the routine RIA quantification of AM, as described in D1, fails to account for the amount of AM bound to its binding protein, factor H. Furthermore, the proposed solution (pretreating the sample with a chaotropic agent to dissociate AM from factor H) is also not disclosed. Therefore, it would not be obvious for a person skilled in the art to introduce this feature in the closest prior art to solve the problem posed. Consequently, the subject-matter of claim 1 is considered to involve an inventive step. The same applies to dependent claims 2-5.

3.2 The subject-matter of claims 6-16 is novel and inventive (Articles 33(2) and 33(3) PCT).
None of the available prior art documents discloses or refers to a complex comprising AM and the newly identified AM binding protein, human complement factor H. Hence, the use of factor H as an AM-binding protein in a method to

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monitor the AM concentrations in a subject is novel and inventive. Therefore, the subject-matter of claim 6 is novel and inventive. The same applies to dependent claims 7-16.

3.3 The subject-matter of claims 17-20 is novel and inventive (Articles 33(2) and 33(3) PCT).

There is no disclosure or reference in any of the available prior art documents of a kit comprising factor H and at least one component to facilitate or detect AM-binding. Therefore, the subject-matter of claim 17 is novel and inventive. The same applies to dependent claims 18-20.

3.4 The subject-matter of claims 21 and 22 is novel and inventive (Articles 33(2) and 33(3) PCT).

Claims 21 and 22 are directed against a method of identifying an antagonist agent which inhibits the complexation of adrenomedullin and factor H. There is no indication in any of the available prior art documents of a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, it would not be obvious for one skilled in the art to identify antagonists which inhibits the formation of the above-mentioned complex. Hence, the subject-matter of claims 21 and 22 is novel and inventive.

3.5 The subject-matter of claim 23 is not inventive (Article 33(3) PCT).

For the assessment of claims of products defined in terms of a process of identification, a product is not rendered novel merely by the fact that it is identified by means of a new process how difficult or time-consuming the technology may be. Claims for products defined in terms of a process of identification are admissible only if the products as such fulfil the requirements of patentability, i.e. that they are novel and inventive.

D4-D6 disclose the existence of an AM-binding protein, such as AMBP-1. Said binding protein would have been identified by the method of claim 21.

Consequently, as it is common practice to provide an antagonist agent in the form of a composition, the subject-matter of claim 23 can not be considered to involve an inventive step (Article 33(3) PCT).

3.6 The subject-matter of claims 24 and 25 is not novel (Articles 33(2) PCT).

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D4 and D5 disclose the existence of an AM-binding protein, such as AMBP-1, which is known to modulate tumor growth. Since said binding protein would have been identified by the method of claim 21, D4 and D5 anticipate the novelty of the subject-matter of claims 24 and 25.

3.7 The subject-matter of claim 26 is novel and inventive (Articles 33(2) and 33(3) PCT).
None of the available prior art documents discloses or refers to a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, the subject-matter of claim 26 is novel and inventive.

3.8 The subject-matter of claims 27 and 28 is novel and inventive (Articles 33(2) and 33(3) PCT).
D3 (page 2626, right column, paragraph antibodies), which is considered to represent the most relevant state of the art, discloses a monoclonal antibody MoAb-G6 which is directed against the PO72 peptide of AM. The problem to be solved by the present invention may therefore be regarded as the provision of an alternative antibody against AM. There is no indication in any of the available prior art documents of a complex comprising AM and the newly identified AM binding protein, human complement factor H. Therefore, it would not be obvious for one skilled in the art to design antibodies against said novel complex. Hence, the subject-matter of claims 27 and 28 is novel and inventive.

3.9 The subject-matter of claims 29-44 and 48-50 is novel and inventive (Articles 33(2) and 33(3) PCT).
The use of the antibody of claim 27 in a pharmaceutical composition (claim 29), in a method of isolating AM/fH complex from a biological sample (claims 30 and 31), in a method of detecting AM/fH complex in a biological sample (claims 32-36), in a kit (claims 37-40), in a method of inhibiting the growth or proliferation of cancer or tumor cells (claims 41-44) and in a method of treating a condition associated with decreased secretion of insulin by pancreatic cells (claims 48-50) is novel and inventive, given that the antibody of claim 27 is novel and inventive.

3.10 The subject-matter of claims 45-47 is novel and inventive (Articles 33(2) and 33(3) PCT).

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There is no disclosure or reference in any of the available prior art documents that a substance which inhibits the complexation of AM and factor H can be used for the treatment of a condition associated with decreased secretion of insulin by pancreatic cells. Therefore, the subject-matter of claims 45-47 is novel and inventive.

3.11 The subject-matter of claim 51 is not novel (Articles 33(2) PCT). D4 discloses the use of factor H in the therapy of autoimmune diseases. Therefore, D4 anticipates the novelty of the subject-matter of claim 51.

3.12 Dependent claim 52 does not contain any features which, in combination with the features of any claim to which it refers, meet the requirements of the PCT in respect of inventive step (Article 33(3) PCT). The feature is merely one of several straightforward possibilities from which the skilled person would select, in accordance with circumstances, without the exercise of inventive skill.

3.13 For the assessment of the present claims 24, 25 and 41-52 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.

4. Section VII

4.1 Page 1 lines 6-9 is irrelevant subject-matter according to Rule 9.1 (iv) PCT.

4.2 The vague and imprecise statement "without departing from the scope and spirit of the invention" in the description (page 58 lines 9-13) implies that the subject-matter for which protection is sought may be different to that defined by the claims, thereby resulting in lack of clarity (Article 6 PCT) when used to interpret them (see also the PCT Guidelines, C-III, 4.3a).

**INTERNATIONAL PRELIMINARY
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4.3 Contrary to the requirements of Rule 5.1(a)(ii) PCT, the relevant background art disclosed in D2, D4 and D5 is not mentioned in the description, nor are these documents identified therein.

5. Section VIII

Claims 24, 25 and 41-52 are directed to methods of treating cancer or diabetes. The application does not meet the requirements of Article 6 PCT because none of these claims enjoy any substantial support in terms of concrete examples in the application as filed. There is no evidence whatsoever in the application as filed that the suggested methods actually yield the desired result.

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
15 March 2001 (15.03.2001)

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C07K 16/00, A61K 38/00

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LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
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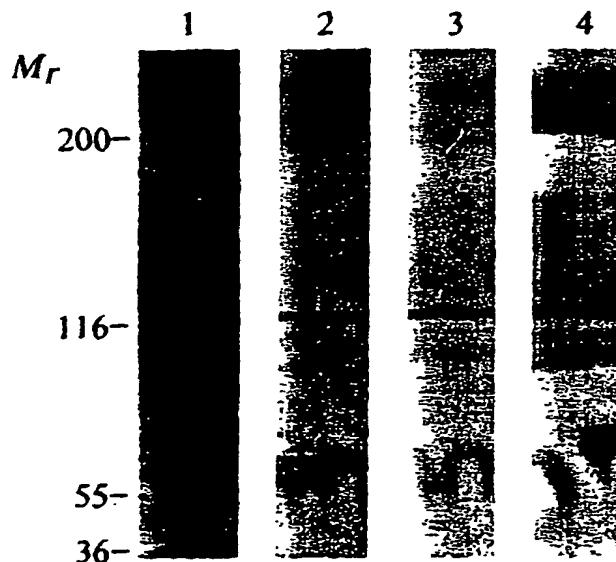
— with international search report

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(88) Date of publication of the international search report:
2 August 2001

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: DETERMINATION OF ADRENOMEDULLIN-BINDING PROTEINS



(57) Abstract: The present invention provides methods for the isolation, identification, and purification of adrenomedullin (AM)-binding proteins. Also, provided are methods for utilizing the purified AM-binding proteins, or functional portions thereof, to diagnose, treat, and monitor AM-related diseases, for example, diseases or disorders associated with abnormally elevated AM levels. In addition, the present invention provides a newly identified complex between AM and a specific AM-binding protein 1 (AMBP-1); which has been isolated and identified herein as factor H (fH). The invention also provides AM/AMBP complexes, particularly AM/fH complexes, and antibodies specifically reactive with these complexes. Further provided are methods for identifying and purifying complexes of AM and an AM binding protein using anti-AM/fH antibodies, and methods for treating conditions such as cancer or diabetes utilizing compositions comprising these antibodies. The present invention additionally provides methods for identifying antagonists agents that inhibit the function of AM, factor H, or the AM/factor H complex. The invention also provides methods for treating conditions such as cancer or diabetes using these antagonist agents.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/24722

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 G01N33/74 C07K16/00 A61K38/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 7 G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, BIOSIS, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MARTINEZ ALFREDO ET AL: "Regulation of insulin secretion and blood glucose metabolism by adrenomedullin." ENDOCRINOLOGY, vol. 137, no. 6, 1996, pages 2626-2632, XP000982198 ISSN: 0013-7227 the whole document ---	1-52
A	WO 97 07214 A (UNSWORTH EDWARD J ;MACRI CHARLES (US); US HEALTH (US); CUTTITTA FR) 27 February 1997 (1997-02-27) the whole document --- -/-	1-52

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

° Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

16 February 2001

Date of mailing of the international search report

26/02/2001

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Pellegrini, P

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/24722

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MARTINEZ ALFREDO ET AL: "Expression of adrenomedullin and its receptor in normal and malignant human skin: A potential pluripotent role in the integument." ENDOCRINOLOGY, vol. 138, no. 12, December 1997 (1997-12), pages 5597-5604, XP000982199 ISSN: 0013-7227 cited in the application the whole document ---	1-52
A	US 4 883 784 A (KANEKO ISAO) 28 November 1989 (1989-11-28) the whole document ---	51,52
P,A	MARTINEZ ALFREDO ET AL: "Adrenomedullin binding protein-1 (AMBP-1) modulates tumor growth by reducing availability of adrenomedullin (AM)." PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page 153 XP000982163 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000 ISSN: 0197-016X abstract ---	1-52
P,A	PIO RUBEN ET AL: "Isolation and characterization of a human plasma adrenomedullin binding protein. Analysis of its expression in human tumor cell lines." PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page 152 XP000982162 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000 ISSN: 0197-016X abstract ---	1-52

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 00/24722

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,A	<p>ELSASSER TED H ET AL: "Adrenomedullin binding protein in the plasma of multiple species: Characterization by radioligand blotting." ENDOCRINOLOGY, vol. 140, no. 10, October 1999 (1999-10), pages 4908-4911, XP000982200 ISSN: 0013-7227 cited in the application the whole document</p> <p>---</p>	1-52
P,A	<p>MARTINEZ A ET AL: "Is adrenomedullin a causal agent in some cases of type 2 diabetes?" PEPTIDES (NEW YORK), vol. 20, no. 12, December 1999 (1999-12), pages 1471-1478, XP000982202 ISSN: 0196-9781 the whole document</p> <p>-----</p>	1-52

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/24722

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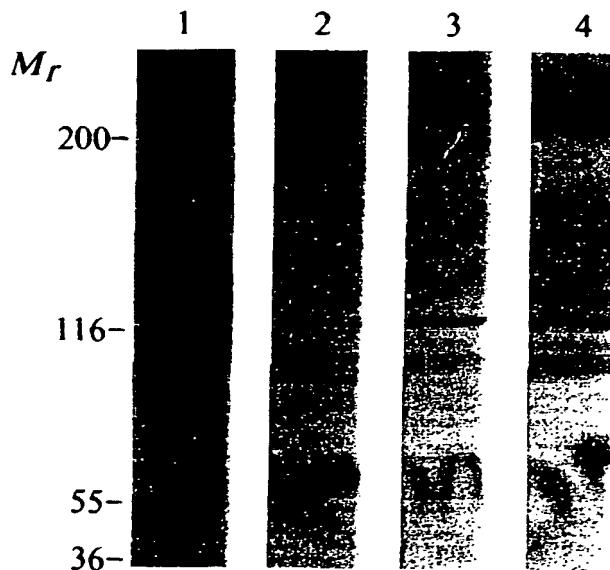
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(54) Title: DETERMINATION OF ADRENOMEDULLIN-BINDING PROTEINS



(57) Abstract: The present invention provides methods for the isolation, identification, and purification of adrenomedullin (AM)-binding proteins. Also, provided are methods for utilizing the purified AM-binding proteins, or functional portions thereof, to diagnose, treat, and monitor AM-related diseases, for example, diseases or disorders associated with abnormally elevated AM levels. In addition, the present invention provides a newly identified complex between AM and a specific AM-binding protein 1 (AMBP-1); which has been isolated and identified herein as factor H (fH). The invention also provides AM/AMBP complexes, particularly AM/fH complexes, and antibodies specifically reactive with these complexes. Further provided are methods for identifying and purifying complexes of AM and an AM binding protein using anti-AM/fH antibodies, and methods for treating conditions such as cancer or diabetes utilizing compositions comprising these antibodies. The present invention additionally provides methods for identifying antagonists agents that inhibit the function of AM, factor H, or the AM/factor H complex. The invention also provides methods for treating conditions such as cancer or diabetes using these antagonist agents.

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**DETERMINATION OF AM-BINDING PROTEINS AND THE ASSOCIATION
OF ADRENOMEDULLIN (AM) THEREWITH**

5

RELATED APPLICATION

This application is related to U.S. Provisional Application Serial Number 60/153,397 filed September 10, 1999, the contents of which are herein incorporated by reference.

10 **FIELD OF THE INVENTION**

The present invention relates to newly developed methods for the detection of adrenomedullin (AM)-binding proteins. The invention further relates to the isolation, identification, and use of such AM-binding proteins, and to compositions and methods including these proteins. The AM-binding 15 proteins can be employed in the diagnosis, treatment and monitoring of AM-related diseases. The present invention also relates to the isolation of AM/AM-binding protein complexes, the generation of antibodies to these complexes, and the use of these antibodies to detect AM/AM-binding protein complexes and treat AM-related diseases. The invention further relates to a 20 newly discovered complex comprising AM and a newly identified AM binding protein, human complement factor H (fH; factor H). The present invention also relates to the identification of antagonists that dissociate the AM/fH complex or inhibit AM/fH complexation, and the use of such antagonists in the prevention and treatment of diseases, including cancers and diabetes.

25 **BACKGROUND OF THE INVENTION**

Adrenomedullin (AM) is a 52 amino acid amidated peptide hormone originally isolated from adrenal tumors. AM is involved in numerous physiological activities, including vasodilation, angiogenesis, mitogenesis, and anti-microbial functions. As a result, AM actions are 30 related to a wide array of disease states such as heart and pulmonary disease, cirrhosis, cancer, diabetes, sepsis, inflammation and preeclampsia

in mammals, including humans. (See, European Patent No. 0 845 036 and U.S. Serial No. 09/011,922, filed February 17, 1998 to F. Cuttiita et al.).

In clinical studies, patients with chronic congestive heart failure show elevated plasma AM, and these concentrations increase in proportion

5 to the severity of the heart failure along with other hormones known to affect the progression of the disease (J. Kato et al., 1996, *J. Clin. Endocrinol. Metab.* **81**:180-3). Similarly, patients with congenital cyanotic heart disease show elevated plasma AM levels and increased AM uptake in pulmonary circulation (M. Yoshibayashi et al, 1999, *Clin. Sci. (Colch.)*, 1999, **96**:543-7).

10 Patients with chronic obstructed pulmonary disease also show significantly raised AM plasma levels (B. Cheung et al., 1997, *Clin. Sci. (Colch.)* **92**:59-62). Accordingly, increased AM (a potent vasodilator) may function as a compensatory mechanism for hypoxaemia (Kato et al., 1996, *supra*; Yoshibayashi et al., 1999, *supra*).

15 Several independent clinical studies have established that patients with liver cirrhosis show elevated circulating AM, and these levels increase with the severity of the illness (E. Fabrega et al., 1997, *Am. J. Gastroenterol.* **92**:1901-4; C.M. Fernandez-Rodriguez et al., 1998, *J. Hepatol.* **29**:250-6; M. Guevara et al., 1998, *Gastroenterology* **114**:336-43; H
20 Kojima et al., 1998, *J. Hepatol.* **28**:840-6). As peripheral vasodilatation has been implicated in the progression of liver cirrhosis, AM may directly participate in the pathogenesis of the disease (Fabrega et al., 1997, *supra*; Fernandez-Rodriguez et al., 1998, *supra*; Guevara et al., 1998, *supra*; Kojima et al., 1998, *supra*).

25 Monoclonal antibodies directed to AM have been shown to inhibit tumor cell growth in a concentration-dependent manner, an effect that is reversed with the addition of exogenous AM (M.J. Miller et al., 1996, *J. Biol. Chem.* **271**:23345-51). AM has also been found to be expressed in numerous and diverse cancer cell lines (M.J. Miller et al., 1996, *supra*), as
30 well as in small and non-small cell lung carcinomas (A. Martínez et al., 1995,

Endocrinology 136:4099-105). In addition, AM binds to specific sites on human malignant melanoma cells and exogenous AM stimulates melanoma cell growth (A. Martínez et al., 1997, *Endocrinology* 138:5597-604). Moreover, cyclic AMP levels in tumor cells increases in the presence of AM 5 (M.J. Miller et al., 1996, *supra*; K. Takahashi et al., 1997, *Peptides* 18:1117-24), indicating that AM may act as a autocrine growth factor to promote neoplastic proliferation (M.J. Miller et al., 1996, *supra*).

It has been demonstrated that AM inhibits insulin secretion in a dose-dependent manner, while neutralizing monoclonal antibodies directed 10 to AM increases insulin release by 5-fold; this effect was reversed by the addition of synthetic AM (A. Martínez, 1996, *Endocrinology* 137:2626-32). Additionally, intravenous injection of AM reduces the levels of insulin in the bloodstream with a concomitant increase in circulating glucose (Martínez, 1996, *supra*). These observations implicate AM as an insulin regulatory 15 factor involved in diabetes and obesity.

In clinical experiments, septic patients showed extremely elevated plasma AM concentrations, and those with acute renal failure had markedly elevated plasma AM levels during the early course of the illness; however, AM levels declined rapidly during the recovery course (Y. Hirata et 20 al., 1996, *J. Clin. Endocrinol. Metab.* 81:1449-53). Similarly, patients with systemic inflammatory response syndrome, pancreatitis, traumatic shock, or severe sepsis show significantly increased plasma levels of AM, and these levels increased in proportion to the severity of illness (S. Ueda et al., 1999, *Am. J. Respir. Crit. Care Med.* 160:132-6). AM levels also correlate with 25 sepsis markers such as the Acute Physiology and Chronic Health Evaluation II score and the peak multiple organ failure score, indicating that AM levels can be used to evaluate the severity of sepsis and can serve as an early predictor of organ failure and outcome (Ueda et al., 1999, *supra*).

The diverse actions of AM are thought to be orchestrated by 30 temporal and/or tissue-specific regulatory factors. The activities of several

other peptide hormones are modulated by binding proteins present in extracellular fluids. For example, one of the most well characterized classes/families of hormone binding proteins are the insulin-like growth factor binding proteins (IGF-BPs). IGF-BPs can direct, enhance, or block 5 the action of IGF-1 on cells by regulating the ability of IGF-1 to bind to cell surface receptors (D.R. Clemmons et al., 1998, *Mol. Cell. Endocrinol.* **140**: 19-24).

The detection, isolation and identification of AM-binding proteins, or families of such proteins, are therefore important goals for the 10 further understanding of AM regulation and function in both normal and disease states in animals, including mammals, preferably humans. Such AM-binding proteins may stabilize or destabilize AM, direct AM to specific sites, modulate AM-binding to its receptor, or otherwise interact with AM to regulate or modulate its activity and/or function. AM-binding proteins may 15 thereby provide a molecular basis for the actions of AM on different tissues, at different times, and in different illnesses and disease states.

Moreover, as a result of the present invention, AM-binding proteins can be used to quantify plasma AM levels in order to diagnose and/or monitor the presence or progression of diseases which are 20 characterized by altered concentrations of circulating AM. AM-binding proteins can also be used to prevent or treat diseases caused or exacerbated by elevated levels of plasma AM by administrating AM-binding proteins in dosages sufficient to bind to AM and thereby block AM activities or interactions with other components.

25 **SUMMARY OF THE INVENTION**

The present invention provides newly developed methods for the detection, isolation and identification of AM-binding proteins, or functional portions thereof, for example, in the form of functional AM polypeptides or peptides. The present invention further provides diagnostics 30 and treatments that utilize the AM-binding proteins, polypeptides, or

peptides. The invention also provides a newly identified complex of AM and an AM-binding protein identified herein as human complement factor H (fH; factor H), (GenBank Accession No. CAA30403). Human complement factor H has been determined to be a marker for urinary bladder carcinomas (R. 5 Heicappell et al., 1999, *Eur. Urol.* 35:81-7). In accordance with the present invention, novel therapeutics which utilize the AM binding protein factor H, or related AM-binding proteins or peptides, are provided for treating cancers, particularly, urinary bladder cancer.

It is an object of the present invention to provide methods of isolating AM-binding proteins, wherein AM is conjugated with a label or marker and incubated with cellular or extracellular lysates. The label or marker may be radioactive or non-radioactive, and is preferably non-radioactive. In accordance with the aspect of the invention related to non-radioactive labels or markers of AM, AM is labeled with fluorescent, 10 chemiluminescent, or immunoreactive molecules, or epitope tags. 15

It is another object of the present invention to provide diagnostic reagents comprising AM-binding proteins or peptides to detect and/or monitor levels of AM in body fluid samples, including cell and tissue lysates and extracts.

It is yet another object of the present invention to provide methods of using AM-binding proteins or peptides as diagnostic reagents for quantifying AM levels, particularly circulating AM levels. Such methods are useful for disease diagnosis, for determining disease severity, and for following the course of treatment for diseases characterized by altered or 20 abnormal AM levels. These diseases include heart and pulmonary diseases, liver cirrhosis, cancers, diabetes, sepsis, inflammation, and other disorders characterized by altered AM plasma concentrations. 25

It is still another object of the present invention to provide improved quantitative assays for detecting AM in serum, utilizing a

chaotropic agent, e.g., sodium thiocyanate to dissociate AM from factor H prior to serum extraction and quantitation of AM.

It is a further object of the present invention to provide quantitative assays for detecting AM, or peptides thereof, using factor H, or
5 related AM-binding polypeptides or peptides, instead of anti-AM antibodies to capture the AM ligand.

It is also a further object of the present invention to provide kits for detecting AM comprising AM-binding proteins, e.g., factor H, or AM-bindable peptides thereof.

10 It is yet a further object of the invention to provide pharmaceutical compositions comprising AM-binding proteins or peptides. In accordance with the present invention, such pharmaceutical compositions are used to treat conditions caused or exacerbated by abnormal, e.g., elevated, levels of plasma AM. These conditions include liver cirrhosis,
15 cancers, diabetes, or other disorders caused or exacerbated by elevated AM plasma concentrations.

It is another object of the present invention to provide a novel complex of AM and the AM-binding protein factor H. The complex is referred to herein as AM/fH.

20 It is also an object of the present invention to provide isolated and substantially purified antibodies that have specific binding affinity or immunoreactivity with an AM/AMBP complex or fragments of the complex, preferably the AM/fH complex or fragments derived therefrom.

It is another object of the present invention to use the above
25 said antibodies in methods to detect the AM/AMBP complex or the AM/fH complex *in vivo* or *in vitro* or treat AM-related conditions, such as cancer or diabetes.

It is still another object of the present invention to provide kits for measuring concentrations of AM/AM-binding protein complex comprising
30 anti-AM/AMBP antibodies, particularly anti-AM/fH antibodies.

It is an additional object of the present invention to provide antagonist agents that inhibit AM, factor H, or AM/fH activity.

It is yet another object of the present invention to provide methods of treating cancer by administering one or more antagonist agents
5 in amounts sufficient to inhibit AM, factor H, or AM/fH activity.

It is also an object of the present invention to provide methods of treating cancer by administering antibodies that specifically bind to the AM/factor H in amounts sufficient to bind to the AM/factor H complex and inhibit AM/fH activity.

10 Further objects and advantages of the present invention will be apparent from the detailed description set forth below.

DESCRIPTION OF THE FIGURES

The appended figures are presented to further describe the invention and to assist in its understanding through clarification of its various
15 aspects.

FIG. 1 illustrates the detection of plasma-derived AM-binding protein 1 (AMBP-1) binding to labeled AM. AMBP-1 has been identified herein as human complement factor H (factor H), based on comparative biochemical and protein determinations and sequence information provided
20 in publicly available databases. Lane 1: AM radiolabeled with ¹²⁵iodine; Lane 2: AM labeled with biotin; Lane 3: AM labeled with fluorescein; Lane 4: AM labeled with dinitrophenol. The band in each lane represents the complex formed by labeled AM binding to AMBP-1 (factor H).

FIG. 2 illustrates the results of a representative competition
25 binding assay to assess the specificity of binding between fluorescein-labeled AM and AMBP-1 (factor H). Lane 1: no competitor added; Lane 2: AM added; Lane 3: insulin-like growth factor (IGF-1) added; Lane 4: proadrenomedullin N-terminal 20 peptide (PAMP) added; Lane 5: insulin added. "f-AM+" indicates fluorescein-labeled AM. The band represents the

AM/AMBP-1 (AM/fH) complex that is unaffected by the presence of competitor peptides.

FIG. 3 illustrates the results of a representative competition binding assay to assess the specificity of binding between fluorescein-labeled AM and AMBP-1 (factor H). Lane 1: no competitor added; Lane 2: full-length AM added; Lane 3: AM₁₋₁₂ added; Lane 4: AM₁₆₋₂₁ added; Lane 5: AM₂₂₋₅₂ added; Lane 6: AM₁₃₋₅₂ added; Lane 7: AM₃₄₋₅₂ added; Lane 8: calcitonin gene-related peptides (CGRP) added; Lane 9: amylin peptides added. "f-AM+" indicates fluorescein-labeled AM. The band represents the AM/AMBP-1 (factor H) complex that is unaffected by presence of competitor peptides.

FIG. 4 illustrates the isolation of AMBP-1. **FIG. 4A** shows fractionation of human plasma by reverse phase HPLC. The dotted line indicates the acetonitrile gradient. **FIG. 4B** shows Coomassie Blue staining of HPLC fractions 47-51. **FIG. 4C** shows ligand blotting of HPLC fractions 47-51.

FIG. 5 illustrates the biochemical characterization of AMBP-1 as human complement factor H. **FIG. 5A** shows Coomassie Blue and glycoprotein staining (GelCode Glycoprotein Staining Kit, Pierce, Rockford, IL) of samples separated by SDS-PAGE. Lane 1: horseradish peroxidase (5 µg), a glycosylated protein used as positive control; Lane 2: soybean trypsin inhibitor (5 µg), a non-glycosylated protein used as negative control; Lane 3: AMBP-1 (2 µg). **FIG. 5B** shows Coomassie Blue staining of samples separated by electrophoresis on a isoelectric focusing gel pH 3-10 (Novex, San Diego, CA). Lane 1: isoelectric focusing markers; Lane 2: AMBP-1 (4 µg). **FIG. 5C** shows Western blot analysis of samples with an anti-factor H antibody. Lane 1: AMBP-1 (100 ng); Lane 2: AMBP-1 (200 ng); Lane 3: commercially available human factor H (50 ng); Lane 4: human plasma (0.2 µl). **FIG. 5D** shows non-radioactive ligand blotting of 1 µg AMBP-1 (Lane 1) and factor H (Lane 2). **FIG. 5E** shows ligand blotting of

AMBP-1 (250 ng) under unreduced (Lane 1) or reduced conditions (Lane 2). FIG. 5F shows binding of fluorescein-labeled AM (50 nM) to a multi-well plate coated with factor H (5 ng/well) is competitively inhibited by increasing concentrations of unlabeled AM (●). In contrast, AM binding to factor H is
5 not affected by PAMP (*) or CGRP (○). Results represent one of two independent experiments. Values represent mean and standard deviation of three determinations. "B/B₀" indicates the ratio of signals generated in the presence/absence of unlabeled competitor.

FIG. 6 illustrates the dissociation of the AM/fH complex. FIG.
10 6A: Purified AMBP-1 (fraction #48) was separated by electrophoresis and transferred to a membrane. After incubation with fluorescein-labeled AM and prior to the final development, the membrane was incubated under various conditions (neutral pH unless indicated). Lane 1: PBS; Lane 2: pH 11.5; Lane 3: pH 2.5; Lane 4: 4M NaCl; Lane 5: 4M NaCl pH 11.5; Lane 6:
15 4M NaCl pH 2.5; Lane 7: 1% SDS; Lane 8: 3M Urea; Lane 9: 3M Guanidine-HCl; Lane 10: 3M sodium thiocyanate (NaSCN); Lane 11: 50% ethylene glycol pH 11.5; Lane 12: 50 % ethylene glycol; Lane 13: 1% β-mercaptoethanol. The band represents the AM/fH complex that remains unaffected by the incubation conditions. FIG. 6B shows dissociation of the
20 AM/fH complex in a multi-well plate assay system. Factor H coated wells were incubated with fluorescein-labeled AM, and prior to the development of the assay, wells were incubated in PBS with 3M NaSCN pH 7.4 for various time periods. Values represent the mean and standard deviation of six determinations. B/B₀ represents the percentage of total binding.

25 FIG. 7 illustrates Western blot analysis of factor H and AM after C18 extraction. FIG. 7A shows human plasma (1 ml) processed with a Sep Pak C18 cartridge and analyzed by Western blotting with anti-factor H antibodies. Lane 1: commercially available human factor H (10 ng); Lane 2: whole human serum (0.5 μl); Lane 3: unbound fraction (1 μl). Lane 4, bound
30 fraction (1 μl). FIG. 7B shows human plasma (1 ml) processed with a Sep

Pak C18 cartridge and analyzed by Western blotting following immunoprecipitation with anti-AM antibodies. Lane 1: synthetic AM (1 ng); Lane 2: fraction immunoprecipitated with normal rabbit serum (30 µl); Lane 3: fraction immunoprecipitated with rabbit anti-AM antibody (30 µl).

5 **FIG. 8** illustrates competitive binding curves generated by human plasma in the AM radioimmunoassay. The dilution curves of plasma (4 ml) extracted following the standard protocol (+) or the NaSCN modification (•) were compared with the standard curve of synthetic AM (o). B/B₀ represents the ratio of radioactivity bound to that bound in the absence 10 of added standard. The scale bar over the curves represents the different plasma dilutions.

15 **FIG. 9** illustrates the effect of factor H on AM activity. **FIG. 9A** shows the effect of factor H in AM mediated cAMP induction. Cyclic AMP was measured after incubating Rat-2 cells with AM and increasing concentrations of factor H. Incubations with AM and 100 or 200 nM factor H produced a significant increase in cAMP as compared to the levels obtained with AM alone ($p<0.01^{**}$ and $p<0.001^{***}$, respectively). Incubation with factor H alone did not affect cAMP levels. Values represent mean and standard deviation of four independent determinations. **FIG. 9B** shows the 20 antimicrobial effect of AM (•), factor H (*), or AM in the presence of 50 µg/ml of factor H (o). Results are expressed in inhibition units (10 units correspond to 1 mm of diameter in the inhibition halo). MIC values were estimated by performing a linear regression and determining the x-intercepts. The MIC value for AM was 18.4 ± 1.3 µg/ml, and increased to 25 35.4 ± 1.1 µg/ml upon the addition of factor H ($p<0.001$). Values represent mean and standard deviation of the mean of eight determinations.

25 **FIG. 10** illustrates the effect of AM on factor H activity. C3b (104 kDa α' chain and 71 kDa β chain) was incubated 24 hr at 37°C with factor H, factor I, and various peptides. Cleavage of the C3b α' chain produced three bands with molecular weights 68 kD, 43 kD, and 42 kD.

FIG. 10A shows the effect of various AM concentrations on the cofactor activity of factor H. **FIG. 10B** shows the effect of AM on factor H activity compared to the effect of the structurally related peptide CGRP and the gene-related peptide PAMP on factor H. Each figure shows a
5 representative example of three different experiments.

FIG. 11 illustrates Northern blot analysis of factor H (4.3 kb) and factor H-like (FHL-1, 1.8 kb) message expression in human tumor cell lines. **FIG. 11A** shows the results of Northern blot analysis using the factor H probe. **FIG. 11B** shows ethidium bromide staining to show the amount of
10 total RNA loaded in each well. "CA" indicates carcinoma; "BAC" indicates bronchioloalveolar carcinoma; "vSCLC" indicates variant small cell lung cancer; "Adeno CA" indicates adenocarcinoma; "Squamous CA" indicates squamous cell carcinoma.

FIG. 12 illustrates immunohistochemical labeling of factor H in
15 rat pancreas. Anti-factor H antibodies localized factor H to cells in the islets of Langerhans (**FIG. 12A**). This localization showed a granular pattern at higher magnification (**FIG. 12B**). Anti-factor H antibodies also localized factor H to some exocrine acini (**FIG. 12C**). Affinity purified anti-factor H antibodies showed the same pattern of staining (**FIG. 12D**). Arrows indicate
20 regions of factor H localization.

FIG. 13 illustrates immunofluorescence labeling of insulin (**FIG. 13A, 13E, and 13I**), factor H (**FIG. 13B, 13F, and 13J**), and either glucagon (**FIG. 13C**), somatostatin (**FIG. 13G**), or pancreatic peptide (**FIG. 13K**) in rat pancreas. The fourth column (**FIG. 13D, 13H, and 13L**) represents an
25 composite of the triple label. In all cases factor H (FH) colocalizes with insulin (INS) in the central β cells, and factor H is absent from the peripheral cells which produce somatostatin (SST), glucagon (GLUC), or pancreatic polypeptide (PP). Samples were analyzed by confocal microscopy.

FIG. 14 illustrates immunogold labeling of factor H (small gold
30 particles, 10 nm in diameter) and insulin (large particles, 20 nm) in rat

pancreas. Low magnification images (FIG. 14A) show that factor H is localized to the characteristic secretory granules of the β cells, and factor H is absent from the surrounding endocrine cell types. High magnification images (FIG. 14B) show that factor H is predominantly localized to the 5 lucent haloes, and insulin is predominantly localized to the dense granule cores. In addition, factor H is present in the secretory granules of some exocrine cells (FIG. 14C). Samples were analyzed by electron microscopy.

FIG. 15 illustrates RT-PCR analysis to determine factor H expression in mouse liver and pancreas. FIG. 15A shows the factor H PCR 10 product (839 bp) amplified from mouse liver and pancreas cells. The PCR product was sequenced to determine that it contained mouse factor H sequence. RT-PCR results were confirmed by Southern blot analysis (FIG. 15B).

FIG. 16 illustrates the effect of factor H on insulin secretion. 15 Incubated of islets with AM and/or factor H resulted in dose-dependent decreases in insulin secretion (FIG. 16A) and concomitant elevations on cAMP in the (FIG. 16B). A representative experiment is shown. Values for each treatment represent the mean and standard deviation of three independent wells. Insulin release is expressed as the ratio between the 20 contents in the high glucose medium divided by the amount in the low glucose one to allow for variations in the number of secreting cells and/or their secretory efficiency. All values were statistically different ($p<0.05$) from the negative controls (first bar).

FIG. 17 illustrates the AM-binding proteins in various species 25 detected by ligand blot analysis using ^{125}I labeled human AM. The bands represent 120 kD or 140 kD moieties identified in the plasma or serum of the ten species analyzed.

FIG. 18 illustrates decreased levels of AM-binding proteins observed in the plasma of parasitized calves as compared to the plasma of

healthy calves. The mean intensity of the bands was decreased by 67% ($P < 0.03$).

DETAILED DESCRIPTION OF THE INVENTION

One embodiment of the present invention encompasses a
5 ligand blotting assay in which AM as ligand is labeled with a radioactive or non-radioactive molecule, marker, or tag, and used as a probe to identify an AM-binding protein or an AM-binding fragment thereof. Preferred in the present invention are non-radioactively labeled AM ligands.

Labeling of AM

10 Numerous methods may be used for the non-radioactive labeling of AM or related peptides. For example, coupling agents such as aldehydes, carbodiimides, dimaleimides, iminodiacetates, succinimides, aminobenzamidines and related compounds can be used to conjugate AM peptides with fluorescent, chemiluminescent, or chemical labels. Examples
15 of nonradioactive labels that can be used include, but are not limited to, fluorescent labels such as fluorescein and its derivatives, e.g., fluorescein isothiocyanate, rhodamine and its derivatives, dansyl, and umbelliferone; chemiluminescers such as 2,3-dihydrophtalazine-dione, and chemical groups such as dinitrophenol (DNP), digoxigenin, and biotin.

20 AM or peptides derived therefrom can also be tagged with amino acid sequences that carry immunoreactive epitopes. A non-limiting list of suitable epitope tags includes c-myc, haemagglutinin (HA), polyhistidine (6X-HIS), GLU-GLU, and DYKDDDDK (FLAG®) tags. Epitope tags can be added to polypeptides or peptides by a number of established
25 methods. The DNA sequences of epitope tags can be inserted into polypeptide or peptide coding sequences as oligonucleotides or via primers used in PCR amplification. Alternately, polypeptide or peptide coding sequences can be cloned into specific vectors that create fusions with epitope tags; for example, pRSET vectors (Invitrogen Corp., San Diego,
30 CA).

The complete AM peptide, or fragments derived from AM, preferably having the same or equivalent function as AM, can be used. AM peptides that may be useful have been described in patent application U.S. Serial No. 09/011,922, filed February 17, 1998 to F. Cuttitta et al.

5 In another aspect of the present invention, AM can be tagged or marked with radioactive isotopes, such as ^{125}I , ^{135}I , ^{35}S , ^{14}C , or [^3H]-thymidine, as nonlimiting examples. Those having skill in the art know how to label proteins and peptides with radioactive isotopic labels using methods and protocols routinely practiced in the art. For example, AM can be labeled
10 by incorporating ^{14}C or ^{35}S labeled amino acids during protein synthesis in host cells or cell-free expression systems (see below). This radiolabeled AM can then be isolated and assayed for its binding to the AM-receptor to confirm its biological function.

Identification of AM-binding proteins

15 Candidate AM-binding proteins, polypeptides, or peptides derived therefrom, can be identified and analyzed by many well-known methods in the art (see T.E. Creighton, Ed., 1997, *Proteins Structure: A Practical Approach*, IRL Press at Oxford Press, Oxford, England). As used herein, the terms protein and polypeptide are synonymous. Peptides are defined as fragments or portions of proteins or polypeptides, preferably fragments or portions having the same or equivalent function or activity as the complete protein.

25 AM-binding proteins can be obtained from biological samples, such as plasma and body fluid samples of animals, including cells and tissues and lysates or extracts derived therefrom. Suitable animal sources of AM-binding proteins include birds, fish, insects, and mammals, including humans. The proteins obtained from these sources can be separated into bands by size fractionation using sodium dodecylsulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred by electroblotting, for example,
30 onto a suitable solid-phase matrix, support, or membrane (e.g., glass or

polymer beads or nylon reinforced nitrocellulose or polyvinylidene fluoride). The support can then be incubated with labeled AM. Proteins/polypeptides that correspond to bands that exhibit specific binding with labeled AM are then identified, isolated/purified, and analyzed by amino acid analysis and

5 Edman degradation to determine the amino acid sequence of peptides derived therefrom.

Sequences determined from Edman degradation can be compared with subject sequences in available databases such as, without limitation, GenBank, SwissProt, BLOCKS, and Pima II. These databases,

10 which contain previously identified and annotated sequences, may be searched for the full-length polypeptide and gene sequence using, for example, Basic Local Alignment Search Tool (BLAST; S.F. Altschul, 1993, *J. Mol. Evol.* 36:290-300; S.F. Altschul et al., 1990, *J. Mol. Biol.* 215:403-10).

In cases where the full-length sequences of AM-binding

15 proteins are not available, extended or overlapping partial clones may be obtained by techniques conventionally known and practiced in the art. Non-limiting examples of such techniques include hybridization to plasmid or phage libraries of genomic DNA or cDNA; PCR from the same libraries using AM-binding protein primer pairs; or hybridization or PCR directly to

20 genomic DNA or cDNA. These clones may then be sequenced and assembled into full-length genes using the fragment sequence alignment program (PHRAP; Nickerson et al., 1997, *Nucleic Acids Res.* 25:2745-2751).

Purification of AM-binding proteins

25 Isolated AM-binding proteins or peptides may be used in diagnostics and treatments according to the present invention. An isolated protein or peptide as used herein refers to a component that is removed from its original environment (for example, its natural environment if it is naturally occurring). An isolated protein or peptide contains less than about

30 50%, preferably less than about 25%, and most preferably less than about

10%, of the components with which it was originally associated. Preferably, an isolated polypeptide or peptide is at least about 80-90% pure, more preferably at least about 90-100% pure.

Both naturally occurring and recombinant forms of the AM-binding proteins or peptides can be used. Methods for directly isolating and purifying polypeptides from natural sources such as cellular or extracellular lysates are well-known in the art (see E.L.V. Harris and S. Angal, Eds., 1989, *Protein Purification Methods: A Practical Approach*, IRL Press, Oxford, England). Such methods include, without limitation, preparative disc-gel electrophoresis, isoelectric focusing, high-performance liquid chromatography (HPLC), reversed-phase HPLC, gel filtration, ion exchange and partition chromatography, and countercurrent distribution, and combinations thereof. Naturally occurring polypeptides can be purified from many possible sources, for example, plasma, body cells and tissues, or body fluids.

To produce recombinant AM-binding proteins or peptides, DNA sequences encoding the AM-binding proteins or peptides are cloned into a suitable vector for expression in intact host cells or in cell-free translation systems (see J. Sambrook et al., 1989, *Molecular Cloning: A Laboratory Manual*, 2nd Edition, Cold Spring Harbor Press, Cold Spring Harbor, NY). Prokaryotic and eukaryotic vectors and host cells may be employed. The particular choice of vector/host/translation system is not critical to the practice of the invention. DNA sequences can be optimized, if desired, for more efficient expression in a given host organism. For example, codons can be altered to conform to the preferred codon usage in a given host cell or cell-free translation system using techniques routinely practiced in the art.

Cloning vectors include, but are not limited to pUC, pBluescript (Stratagene, La Jolla, CA), pET (Novagen, Inc., Madison, WI) and pREP (Invitrogen Corp.) plasmids. Vectors can contain one or more replication and inheritance systems for cloning or expression, one or more markers for

selection in the host, e.g. antibiotic resistance, and one or more expression cassettes. The inserted coding sequences can be synthesized by standard methods, isolated from natural sources, or prepared as hybrids. Ligation of the coding sequences to transcriptional regulatory elements (e.g., 5 promoters, enhancers, and/or insulators) and/or to other amino acid encoding sequences can be carried out using established methods.

For some purposes, it may be preferable to produce peptides or polypeptides in a recombinant system wherein the peptides or polypeptides carry additional sequence tags to facilitate purification. Such 10 markers include epitope tags (described above) and protein tags, for example, glutathione-S-transferase (GST), green fluorescent protein (GFP), and maltose binding protein (MBP). Protein tags are attached to peptides or polypeptides by several well-known methods. As a non-limiting example, the coding sequence of a polypeptide or peptide can be cloned into a vector 15 that creates a fusion between the polypeptide or peptide and a protein tag of interest. Suitable vectors include, without limitation, the exemplary plasmids, pGEX (Amersham Pharmacia Biotech, Inc., Piscataway, NJ), pEGFP (CLONETECH Laboratories, Inc., Palo Alto, CA), and pMALTM (New England BioLabs, Inc., Beverly, MA). The tagged polypeptide or peptide can 20 then be purified from a crude lysate of the translation system or host cell by chromatography on an appropriate solid-phase matrix.

Suitable cell-free expression systems for use in accordance with the present invention include rabbit reticulocyte lysate, wheat germ extract, canine pancreatic microsomal membranes, *E. coli* S30 extract, and 25 coupled transcription/translation systems (Promega Corp., Madison, WI). These systems allow the expression of recombinant polypeptides or peptides upon the addition of cloning vectors, DNA fragments, or RNA sequences containing coding regions and appropriate promoter elements.

Host cells for recombinant cloning vectors include bacterial, 30 archebacterial, fungal, plant, insect and animal cells, especially mammalian

cells. Of particular interest are *E. coli*, *Bacillus subtilis*, *Staphylococcus aureus*, *Saccharomyces cerevisiae*, *Schizosaccharomyces pombe*, *Neurospora crassa*, SF9, C129, 293, NIH 3T3, CHO, COS, and HeLa cells. Such cells can be transformed, transfected, or transduced, as appropriate,

5 by any suitable method including electroporation, CaCl_2 -, LiCl -, LiAc/PEG -, spheroplasting-, Ca-Phosphate, DEAE-dextran, liposome-mediated DNA uptake, injection, microinjection, microprojectile bombardment, or other established methods.

Antibody-based methods may also be used to purify natural or
10 recombinantly produced AM-binding proteins or peptides. Antibodies that recognize these polypeptides, or peptides derived therefrom, can be produced and isolated using methods known and practiced in the art. AM-binding polypeptides or peptides can then be purified from a crude lysate by chromatography on antibody-conjugated solid-phase matrices (see E.
15 Harlow and D. Lane, 1999, *Using Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY). Other purification methods known and used in the art may also be employed.

The AM-binding protein Human Complement Factor H (fH)

In a preferred embodiment of the present invention, the AM-
20 binding protein is human complement factor H. Factor H was originally identified as an important modulator of the complement cascade of the immune system (see M.P. Dierich, 1988, *Mol. Immunol.* **25**:1043-51). Specifically, factor H forms a complex with factor I, and the fH-fI complex selectively degrades C3b, inactivates the C3 convertase, and competes for
25 factor B binding to C3b (P.F. Zipfel, et al., 1999, *Immunopharmacol.* **42**:53-60). In addition, factor H can bind to the integrin receptor (CD11b/CD18) expressed in a variety of human tissues, and bind to certain surface glycosaminoglycans present on both prokaryotic and eukaryotic cells (P.F. Zipfel, et al., 1999, *Immunopharmacol.* **42**:53-60).

The full length cDNA (4.3 kb) for human factor H has been cloned and sequenced and found to undergo alternative gene splicing to give rise to a shorter message (1.8 kb) (P.F. Zipfel et al., 1999, *Mol. Immunol.* 36:241-8). This smaller message is translated into a truncated factor H molecule (43 kDa), denoted as factor H-like protein (FHL-1), that shares the complement regulatory functions of factor H (P.F. Zipfel et al., 1999, *Mol. Immunol.* 36:241-8). Structurally, factor H contains twenty short consensus repeats (SCRs), while FHL-1 contains seven SCRs. According to conventional nomenclature, factor H is represented as SCR 1-20 with FHL-1 being SCR 1-7 (P.F. Zipfel, et al., 1999, *Immunopharmacol.* 42:53-60). Notably, both factor H and factor H-like proteins have been used as markers for urinary bladder carcinomas (R. Heicappell et al., 1999, *Eur. Urol.* 35:81-7).

In accordance with the AM-binding protein isolation and identification methods described herein, the present inventors have demonstrated that factor H is an AM-binding protein isolated from human plasma. Specifically, a non-radioactive ligand blotting technique (Examples 1 and 2; FIG. 1) and HPLC/SDS-PAGE (Example 4; FIGS. 4A-4C) purification techniques allowed the isolation of the AMBP-1 protein. The purified protein was subjected to total amino acid analysis, N-terminal amino acid sequence, and mass spectrometry (Example 5; Table 1). In addition, purified AMBP-1 was compared with factor H, based on apparent molecular weight, glycosylation, AM-binding, and recognition by anti-factor H antibodies (Example 6; FIGS. 5A, 5C, and 5D). The results identified AMBP-1 as human complement factor H.

Additional experiments as described hereinbelow were conducted to characterize the AM/fH interaction (Example 7). These experiments showed that the AM/fH complex was not dissociated in acidic or high salt conditions (FIG. 6A). In contrast, the AM/fH complex was dissociated in the presence of chaotropic agents, such as NaSCN (FIG. 6A).

The dissociation of the AM/fH complex by NaSCN was confirmed using a multi-well AM-binding assay, 96-well plate (Example 7). The displacement curve suggested the presence of two dissociation mechanisms, as shown by an initial rapid dissociation phase, followed by a slower dissociation phase
5 (FIG. 6B).

Notably, the experiments described herein have shown that the AM/fH interaction interferes with the established methodology for quantification of circulating AM (Example 8). Thus, the routine radioimmunoassay (RIA) quantification of AM fails to account for the amount
10 of AM bound to its binding protein, factor H. The inventors' studies have indicated that the standard C18 reverse-phase separation technique used to prepare plasma for RIA analysis effectively eliminates AMBP-1 (factor H) from the extract (T. H. Elsasser et al., 1999, *Endocrinology* 140:4908-4911; FIG. 7A). Further studies have demonstrated that AM is detected in both
15 the unbound and the bound fractions following C18 reverse-phase separation (FIG. 7B), confirming that the traditional procedure does not recover the total amount of AM present in plasma. In addition, the inventors have performed a modified RIA analysis utilizing plasma pre-treated with a chaotropic agent, NaSCN, to dissociate AM/fH complexes in the plasma
20 prior to extraction. The modified protocol detected AM levels that were twofold higher than those obtained with the standard technique (FIG. 8).

The inventors' experiments have further shown that factor H modulates AM activity, and AM modulates factor H activity. In particular, factor H enhances AM-mediated induction of cAMP in fibroblasts (Example
25 9; FIG. 9A), but suppresses the bactericidal activity of AM against *E. coli* (Example 9; FIG. 9B). Additionally, AM enhances fH/fI-mediated cleavage of C3b (Example 9; FIGS. 10A and 10B). Thus, the interaction of AM and factor H produces wide-ranging effects on both AM and factor H function.

In addition, experiments described in detail herein have shown
30 that factor H and FHL-1 are highly expressed in non-small cell lung cancer

cells, adenocarcinoma cells, and squamous cell carcinoma (Example 10; FIG. 11). Factor H and FHL-1 are also expressed in other solid tumor cells, such as breast, colon, ovary, and prostate carcinoma cells (FIG. 11). These results are consistent with recent findings, which have determined that factor

5 H is expressed in a variety of human cancer cells (i.e., bladder cancer, breast cancer, and glioblastoma cells), and factor H facilitates cancer cells' resistance to complement-mediated cell lysis (N.S. Fedarko et al., 2000, *J. Biol. Chem.* **275**:16666-16672; S. Junnikkala et al., 2000, *J. Immunol.* **164**: 6075-6081; P.F. Zipfel et al., 1999, *Mol. Immunol.* **36**:241-8).

10 Similarly, the inventors' studies have shown AM is expressed in several cancer cell lines, and monoclonal antibodies directed to AM inhibit tumor or cancer cell growth in a concentration-dependent manner (Martínez et al., 1995, *supra*; M. Garayoa et al., 2000, *Mol. Endocrinol.* **14**(6):848-862; Martínez et al., 1997, *supra*; Miller et al., 1996, *supra*). Moreover, the

15 experiments described herein show that AM potentiates fH/fI-mediated degradation of C3b (Example 9; FIG. 10A and 10B). The sum of these observations indicates that the AM/fH complex may mediate cancer cells' resistance to complement-mediated cell lysis. Accordingly, antagonists that inhibit AM activity, inhibit factor H activity, or prevent the interaction of AM

20 and factor H may act to decrease cancer cells' resistance to cell lysis. Such antagonist agents may thereby be used as cancer therapeutics.

In a separate set of experiments described hereinbelow, the inventors have used immunohistochemical labeling (Example 11; FIGS. 12A-12D) and multiple immunofluorescence labeling followed by confocal

25 microscopy (Example 12; FIGS. 13B, 13F, and 13J) to demonstrate that factor H is expressed by the β cells of rat pancreatic islets. Double immunogold staining under the electron microscope also showed colocalization of insulin and factor H within the same secretory granules (Example 13; FIG. 14B). Factor H was localized to electron-lucent haloes,

30 whereas insulin was localized predominantly to electron-dense cores (FIG.

14B). The presence of factor H mRNA in the pancreas was confirmed by RT-PCR analysis (Example 14; FIG. 15A).

The inventors have previously determined that AM acts to reduce insulin secretion in the pancreas (A. Martínez, 1996, *Endocrinology* 137:2626-32). Thus, insulin secretion assays described herein have been used to investigate the role of factor H in pancreatic function (Example 15). Such assays have demonstrated that in the presence of AM, factor H induces a further reduction in insulin secretion with a concomitant elevation of cAMP (FIGS. 16A and 16B). Thus, antagonists that inhibit AM activity, inhibit factor H activity, or prevent the interaction of AM and factor H may act to increase insulin production. Such antagonist agents may thereby be used as therapeutics for conditions associated with reduced secretion of insulin (e.g., certain forms of type 2 diabetes and pheochromocytoma).

As indicated above, factor H can act to suppress complement-mediated cell lysis. In particular, factor H has been shown facilitate the resistance of bacterial cells (C. Neeleman et al., 1999, *Infect. Immun.* 67:4517-4524) and tumor cells (N.S. Fedarko et al., 2000, *J. Biol. Chem.* (in press)) to complement-mediated lysis. It is therefore possible that factor H normally protects β cells from complement-mediated lysis, and inhibition or reduced expression of factor H in β cells results in the autoimmune destruction of the cells. It is further possible that AM assists factor H in providing β cells with protection against complement-mediated lysis (see FIGS. 10A and 10B). In this way, factor H, the AM/fH complex, or functional fragments thereof may be used to treat conditions associated with autoimmune responses against β cells (e.g., type 1 diabetes).

The inventors have further demonstrated that plasma proteins from several species can specifically bind AM (T.H. Elsasser et al., 1999, *Endocrinology* 140:4908-4911; Example 16; FIG. 17). These AM-binding proteins have been identified by a radioligand blotting technique adapted from P. Hossenlopp et al. (1986, *Anal. Biochem.* 154:138-143). An AM-

binding protein of molecular weight (M_r) 120 kD (under non-reducing conditions) has been observed in the plasma from most of the species analyzed, including humans. It is noted that the human 120 kD AM-binding protein corresponds to AMBP-1 (factor H), described in detail herein.

5 Interestingly, an additional band of M_r 140 kD has been observed in the plasma from ruminant species (i.e., calf, goat, and sheep). This band may represent a different protein or a different glycosylation pattern of the 120 kD protein. From these experiments, it is concluded that AM-binding proteins in animal and avian species may interact with AM to form AM/AMBP

10 complexes similar to the AM/fH complex described herein.

Antibodies to AM-binding proteins

The present invention provides antibodies that specifically recognize AM/AMBP complexes or fragments of these complexes, preferably AM/fH complexes or fragments derived therefrom. The present

15 invention also provides assay methods comprising these antibodies to identify and/or distinguish among AM/AMBP complexes, preferably the AM/fH complex, *in vivo* or *in vitro*. For example, the methods involve the use of the antibodies, or labeled antibodies, that have been bound to a solid support or matrix in assays, e.g., immunoassays, to bind to the AM/AMBP

20 complex, or a complex-specific fragment thereof.

In accordance with the invention, specific antibodies can be generated that recognize and bind to a specific complex of AM and an AM-binding protein to aid in the identification and detection or isolation of the complex. Preferred are antibodies that recognize and bind to the AM/fH complex, but do not recognize and bind to either AM or factor H alone. One method for utilizing the antibodies according to the present invention to isolate AM/AM binding protein complexes in a biological sample comprises providing a solid support to which are bound antibodies that recognize a specific AM/AM binding protein complex, or a complex-specific fragment

25 thereof, contacting the support with the sample or an aliquot of the sample

30

and eluting the complex that binds to the antibodies adsorbed onto the support. Another method for detecting AM/AM binding protein complexes in a sample comprises incubating the sample with antibodies that specifically recognize and bind to a complex of AM and an AM binding protein, or a
5 complex specific fragment, under conditions that allow the antibodies to bind to the AM/AM binding protein complex and determining the binding of the antibodies to the complex.

As used herein, "antibody" refers to intact molecules as well as fragments thereof, such as Fab, F(ab)₂, and Fv, which are capable of
10 binding an epitopic determinant. Antibodies that bind to the AM/AMBP complex, preferably AM/fH, can be prepared using the isolated AM/AMBP complex, preferably, AM/fH, or fragments containing small peptides specific to the complex as the immunogen or immunizing antigen. Antibodies can also be generated to AM-binding proteins or fragments derived therefrom
15 using the isolated AM-binding protein or a derived fragment, as an immunogen. As will be appreciated by those having skill in the art, the immunogen can be conjugated to a carrier protein, if desired, to increase immunogenicity, particularly, if a small peptide is used. Commonly used carriers that are routinely used chemically coupled to peptides include serum
20 albumins, i.e., bovine, sheep, goat, or fish serum albumin, thyroglobulin, and keyhole limpet hemocyanin. The coupled immunogen-carrier is then used to immunize a recipient animal (e.g., mouse, rat, sheep, goat, or rabbit).

The term "antigenic determinant" refers to that fragment of a molecule (i.e., an epitope) that makes contact with a particular antibody.
25 When the AM/AMBP complex is used to immunize a host animal, numerous regions of the complex may induce the production of antibodies which bind specifically to a given region or three-dimensional structure on the complex; these regions or structures are referred to as antigenic determinants or epitopes. An antigenic determinant may compete with the intact antigen
30 (i.e., the immunogen used to elicit the immune response) for binding to an

antibody. Preferred are those antigenic determinants that are specific for the AM/AMBP complex.

AM/AMBP complex-specific antibodies according to the present invention include polyclonal and monoclonal antibodies. The 5 antibodies can be elicited in an animal host by immunization with AM/AMBP complex-derived immunogenic components or can be formed by *in vitro* immunization (sensitization) of immune cells. The immunogenic components used as immunogens to elicit the production of antibodies can be isolated from plasma, recombinantly produced, or chemically 10 synthesized. The antibodies can also be produced in recombinant systems programmed with appropriate antibody-encoding DNA. Alternatively, the antibodies can be constructed by biochemical reconstitution of purified heavy and light chains. The antibodies include hybrid antibodies, chimeric antibodies, humanized antibodies (see, for example, U.S. Patent No. 15 5,585,089) and univalent antibodies. Also included are Fab fragments, Fab' and F(ab)₂ fragments of antibodies.

AM/AMBP complex as immunogens can be obtained using techniques as described above. For example, AM/AMBP complexes can be isolated and purified by excising the complexes from gels, particularly SDS- 20 PAGE gels. Alternatively, recombinant AM and AM-binding proteins or related fragments can be co-expressed in a host cell or cell-free expression system and co-purified.

The immunogenic components of the AM/AM binding protein complex according to the invention are useful as antigens for preparing 25 antibodies by standard methods. These antibodies, whether polyclonal or monoclonal, can be used, for example in an immobilized form bound to a solid support by well-known methods, to purify the immunogenic components, specifically, AM/AMBP complexes such as AM/fH, by immunoaffinity chromatography. In addition, the AM/AM binding protein

complex can be used to screen antibodies, particularly monoclonal antibodies, that are generated as described below.

Hybridomas that produce monoclonal antibodies against the immunogenic components of the invention can be produced by well-known techniques. Hybridomas can be produced by the fusion of an immortalized cell line with a B-lymphocyte that produces the desired antibody. Alternatively, non-fusion techniques for generating immortal antibody-producing cell lines are possible, and are within the purview of the present invention (see Casali et al., 1986, *Science*, **234**:476). Immortalized cell lines are typically transformed mammalian cells, particularly myeloma cells of rodent, bovine, and human origin. Most frequently, rat or mouse myeloma cell lines are employed as a matter of convenience and availability.

Standard procedures can be used to select hybridomas, such as HAT (hypoxanthine-aminopterin-thymidine) selection. Hybridomas that secrete desired monoclonal antibodies can be selected by assaying the cells' culture medium by standard immunoassays, such as immunoblotting, ELISA (enzyme-linked immunosorbent assay; E. Engvall et al., 1971, *Immunochemistry*, **8**:871-4; and D.J. Reen, 1994, *Methods Mol. Biol.* **32**:461-6), RIA, or comparable assays. Antibodies can be recovered from the medium using standard protein purification techniques (see Tijssen, 1985, *Practice and Theory of Enzyme Immunoassays*, Elsevier, Amsterdam).

Agonists/Antagonists of AM, AM-binding proteins, or factor H

Agonists/antagonists that modulate AM activity, AM-binding protein (e.g., factor H) activity, or affect AM/AMBP (e.g., AM/fH) interaction may be employed as therapeutics. For example, antagonists that inhibit AM or factor H activity, or block AM/fH interaction may be formulated into pharmaceutical compositions that are used to inhibit the growth or proliferation of cancer cells, in particular, bladder cancer, breast cancer, and glioblastoma cells. Alternatively, agonists that increase AM activity and

antagonists that decrease factor H activity or prevent AM/fH interaction can be formulated into pharmaceutical compositions and used to treat microbial infection, in particular, bacterial infection.

Modulators may comprise nucleic acids, oligonucleotides;

5 polypeptides, peptides; oligosaccharides; lipids; antibodies, or derivatives or fragments of any of the foregoing, or other organic or inorganic molecules. Modulators can be identified using methods well-known in the art, such as, for example, by screening natural product libraries, fermentation libraries (encompassing plants and microorganisms), combinatorial libraries,

10 compound files, and synthetic compound libraries for compounds that have the ability to bind to, and/or alter the function of AM or AM-binding protein (e.g., factor H). In particular, synthetic compound libraries can be obtained from commercial sources (e.g., Maybridge Chemical Co. (Trevillet, Cornwall, UK); Comgenex (Princeton, NJ); Brandon Associates (Merrimack, NH); and

15 Microsource (New Milford, CT). In addition, a rare chemical library can be obtained from Aldrich Chemical Company, Inc. (Milwaukee, WI). Alternatively, libraries of natural compounds in the form of bacterial, fungal, plant and animal extracts are available from, for example, Pan Laboratories (Bothell, WA), or are readily producible. Additionally, natural and

20 synthetically produced libraries and compounds are readily modified through conventional chemical, physical, and biochemical means (Blondelle et al., 1996, *Trends in Biotech.* 14:60).

In one embodiment of the present invention, modulators may be identified by screening test molecules against AM or AM-binding protein

25 (e.g.,) nucleic acid or amino acid sequences in high-throughput assays. Such assays include without limitation genetic, biochemical, and ligand binding assays. Several methods of automated assays have been developed to permit screening of tens of thousands of compounds in a short period of time. These methods are particularly preferred. The use of high-

30 throughput screening assays to test for modulators is greatly facilitated by

the availability of large amounts of purified nucleic acid or amino acids sequences, as provided by the invention.

Ligand-binding assays can be used to detect binding of test compounds to particular sequences. The detection may involve direct measurement of binding. Alternatively, indirect indications of binding may involve stabilization of protein structure or disruption of a biological function. Non-limiting examples of useful ligand-binding assays are detailed below.

A useful method for the detection and isolation of binding proteins or peptides is the Biomolecular Interaction Assay (BIAcore) system (Pharmacia Biosensor, LKB Pharmacia, Sweden). The BIAcore system uses affinity purified anti-GST antibodies to immobilize GST-fusion proteins onto a sensor chip. A protein or peptide of interest is coated onto a chip and test compounds are passed over the chip. Binding is detected by a change in the surface plasmon resonance, which is an optical phenomenon that detects changes in refractive indices.

Alternatively, binding proteins can be identified by scintillation proximity assays (SPAs, described in U.S. Patent No. 4,568,649). In a modified version of this assay, a tagged protein is attached to SPA beads, and test compounds are added. The tagged protein is then subjected to mild denaturing conditions (such as, e.g., heat, exposure to SDS, etc.) and a purified labeled chaperonin is added. If a test compound has bound to a tagged protein, the labeled chaperonin will not bind; conversely, if no test compound has bound, the chaperonin will bind.

In another approach, binding proteins or peptides can be identified using the binding assay described in Fodor et al., 1991, *Science* 251:767-773). Binding proteins can also be identified by *in vitro* mitochondrial-targeting assays (MTAs, based on Hurt et al., 1985, *EMBO J.* 4:2061-2068; Eilers and Schatz, *Nature*, 1986, 322:228-231). In accordance with MTAs, expression vectors are constructed to comprise a coding sequence for a polypeptide or peptide fused with a coding sequence

for a mitochondrial localization signal. The resulting fusion proteins are produced and tested for importation into isolated mitochondria in the presence or absence of a test compound. It is predicted that a test compound that binds to a fusion protein will inhibit mitochondrial import.

5 In addition, binding proteins can be identified by the yeast two-hybrid system (Fields and Song, 1989, *Nature* 340:245-246; U.S. Patent No. 5,283,173). The two-hybrid system assays the reconstitution of transcription activity through the association of a DNA-binding domain and a transcription activation domain of a transcriptional activator via protein-protein
10 interactions. The yeast GAL4 transcriptional activator may be used in this system, although other transcription factors have been used. In brief, the GAL4 DNA-binding domain and the transcription activation domain are expressed, separately, as fusions to potential interacting polypeptides or peptides. If the two, coexpressed fusion proteins are targeted to the nucleus
15 and interact, activation of a reporter gene (e.g. LacZ) produces a detectable phenotype. Related *in vivo* methods such as the three-hybrid (Licitra and Liu, 1996, *Proc. Natl. Acad. Sci. USA* 93:12817-12821), and reverse two-hybrid (Vidal *et al.*, 1996, *Proc. Natl. Acad. Sci. USA* 93:10315-10320) systems may serve as alternative approaches.

20 As not all sequences are suitable for ligand binding assays, other types of assays, e.g., cell-free biochemical assays, are also contemplated for use. It will be appreciated by those skilled in the art that different types of assays can be used to detect different types of modulators.
Diagnostics utilizing AM-binding proteins

25 Another embodiment of the present invention is directed to *in vitro* screening and diagnostic methods for detecting or monitoring the presence of AM or AM-binding proteins, preferably factor H or related proteins or peptides, in biological samples, e.g., serum, plasma, or cell and tissue extracts or lysate. For use as screening and diagnostic reagents in
30 the methods, AM, AM-binding proteins or peptides, preferably factor H or

related proteins or peptides, can be labeled by several established methods in the art. Polypeptides or peptides can be conjugated to labels using a number of different coupling agents as described above, and suitable labels include, without limitation, enzyme-based, fluorescent, chemiluminescent, radioactive, or dye molecules for use in assays that involve such labeled AM-binding proteins. Suitable assays that amplify the signals from the target or probe are well-known, e.g., biotin and avidin assays, and ELISAs can be used with the labeled AM-binding proteins to detect or monitor AM levels.

To provide a basis for the diagnosis of AM-related diseases associated with abnormal levels of AM, a normal or standard profile for AM expression may be established. This can be accomplished by incubating body fluids, tissue preparations, or cell lysates or extracts, for example, obtained from normal subjects, either animal or human, under conditions suitable for binding with labeled AM-binding proteins or peptides. Standard binding levels of AM to AM-binding proteins or peptides may be quantified by comparing the values obtained from normal subjects with those obtained from a standard or control sample in which a known amount of substantially purified AM is used. Standard values obtained from normal samples can be compared with values obtained from samples from subjects who are symptomatic for illness. A significant deviation between standard and test or experimental values is then used to determine the presence and severity of a disorder or disease involving abnormal or elevated levels of AM, particularly, circulating AM in plasma.

Following a diagnosis of a disorder or disease, and initiation of a treatment, binding assays can be repeated on a regular basis to evaluate whether the levels of AM, e.g., plasma AM, in the patient changes, or begins to approximate that which is observed in a normal individual. The results obtained from successive assays can be used to show the efficacy of treatment over a period of time, ranging from several days to months.

Additionally, AM-binding proteins or peptides may be included as reagents in AM detection or diagnostic kits. Suitable kits may include one or more of the following components:

- 5 i) one or more AM-binding proteins or peptides, preferably factor H or related functional peptides thereof; the included AM-binding proteins or peptides may be pre-labeled; alternatively, the AM-binding proteins or peptides may be unlabelled and the ingredients for labeling may be included in the kit in separate containers; and
- 10 ii) reaction components such as buffers or reagents to mediate or measure AM-binding. The kit may also contain other suitably packaged reagents and materials needed for the binding protocol, including, for example, solid-phase matrices, standards, and instructions for conducting the test.

A further embodiment of the present invention is directed to *in vitro* screening and diagnostic methods for detecting or monitoring the presence of the AM/fH complex in biological samples, e.g., serum, plasma, or cell and tissue extracts or lysate. For use as screening and diagnostic reagents in the methods, antibodies that specifically recognize the AM/fH complex can be utilized according to well-established methods in the art.

20 Such methods comprise the steps of:

- 25 i) contacting a sample suspected to contain altered levels of the AM/fH complex with an antibody specific for the complex under conditions in which a stable AM/fH-antibody association can form between the antibody and AM/fH complex in the sample; and
- ii) detecting any AM/fH-antibody association formed in step (i) using any suitable means known in the art, wherein the amount of the AM/fH-antibody association detected indicates the level of AM/fH complex in the sample.

Many immunoassay formats are known in the art, and the 30 particular format used is determined by the desired application. An

immunoassay may use, for example, a monoclonal antibody directed against a single AM/fH complex epitope or a combination of monoclonal antibodies directed against different epitopes of the AM/fH complex. Protocols may also, for example, use solid supports, or may involve immunoprecipitation.

5 Examples of solid supports that can be used include, without limitation, nitrocellulose (e.g., in membrane or microtiter well form), polyvinyl chloride (e.g., in sheets or microtiter wells), polystyrene latex (e.g., in beads or microtiter plates), polyvinylidene fluoride (known as IMMUNOLON TM), diazotized paper, nylon membranes, activated beads, and protein A beads.

10 For example, Dynatech IMMUNOLON TM 1 or IMMUNOLON TM 2 microtiter plates or 0.23 inch polystyrene beads (Precision Plastic Ball) can be used.

15 Typically, immunoassays use either a labeled antibody or a labeled antigenic component (e.g., that competes with the antigen in the sample for binding to the antibody). Suitable labels include without limitation enzyme-based, fluorescent, chemiluminescent, radioactive, or dye molecules. Assays that amplify the signals from the probe are also known, such as, for example, those that utilize biotin and avidin, and enzyme-labeled immunoassays, such as ELISA assays.

20 Kits suitable for antibody-based diagnostic applications typically include one or more of the following components:

(i) Anti-AM/fH complex antibodies: The antibodies may be pre-labeled; alternatively, the antibody may be unlabeled and the ingredients for labeling may be included in the kit in separate containers, or a secondary, 25 labeled antibody is provided; and

(ii) Reaction components: The kit may also contain other suitably packaged reagents and materials needed for the particular immunoassay protocol, including solid-phase matrices, if applicable, and standards.

The kits may also include instructions for conducting the test. Furthermore, in preferred embodiments, the diagnostic kits are adaptable to high-throughput and/or automated operation.

Treatments utilizing AM-Binding Proteins

5 Another embodiment of the present invention embraces treatments and therapeutic methods that comprise AM-binding proteins or peptides. Because AM-binding proteins or peptides are naturally occurring plasma components, they may be administered to an individual's circulatory system with minimal risk of immunological complications. Pharmaceutical
10 compositions can be produced and employed in treatment protocols according to established methods depending on the disorder or disease to be treated (see, for example, P.D. Mayne, 1996, *Clinical Chemistry in Diagnosis and Treatment*, 6th ed., Oxford University Press, Oxford, England; Gilman et al., eds., 1990, *Goodman and Gilman's: The Pharmacological
15 Basis of Therapeutics*, 8th ed., Pergamon Press; Avis et al., eds., 1993, *Pharmaceutical Dosage Forms: Parenteral Medications*, Dekker, New York; and Lieberman et al., eds., 1990, *Pharmaceutical Dosage Forms: Disperse Systems*, Dekker, New York).

In one embodiment of the present invention, peptides derived
20 from factor H may be used as therapeutics. Such peptides can be produced by recombinant or synthetic techniques and then administered to tumor or cancer cells to inhibit the growth and/or proliferation of these cells. Non-limiting examples of cancer or tumor cells for use with the present invention include urinary bladder, urethral, renal, rectal, colon, small intestine, gastric,
25 esophageal, salivary gland, gallbladder, liver, breast, vaginal, endometrial, ovarian, cervical, prostate, skin, lung, or brain cancer or tumor cells. It is noted that peptide-based treatments would be particularly advantageous, as peptides would not be expected to produce an undesirable immune response in the host, which may be observed during treatments using
30 mouse monoclonal antibodies such as MoAb-G6 (Miller et al., 1996, *supra*).

In an alternative embodiment of the present invention, antibodies specifically bind to the AM/fH complex may be used as therapeutics. For example, anti-AM/fH antibodies can be formulated into a pharmaceutical composition and used to inhibit the growth or proliferation of 5 cancer cells, in particular bladder cancer, breast cancer, and brain cancer cells.

It is noted that antibody-based therapeutics produced from non-human sources can cause an undesired immune response in human subjects. To minimize this problem, chimeric antibody derivatives can be 10 produced. Chimeric antibodies combine a non-human animal variable region with a human constant region. Chimeric antibodies can be constructed according to methods known in the art (see Morrison et al., 1985, *Proc. Natl. Acad. Sci. USA* 81:6851; Takeda et al., 1985, *Nature* 314:452; U.S. Patent No. 4,816,567; U.S. Patent No. 4,816,397; European 15 Patent Publication EP 171496; EP 0173494; United Kingdom Patent GB 2177096B). In addition, antibodies can be further "humanized" by any of the techniques known in the art, (e.g., Teng et al., 1983, *Proc. Natl. Acad. Sci. USA* 80:7308-7312; Kozbor et al., 1983, *Immunology Today* 4: 7279; Olsson et al., 1982, *Meth. Enzymol.* 92:3-16; International Patent Application 20 WO92/06193; EP 0239400). Humanized antibodies can be also be obtained from commercial sources (e.g., Scotgen Limited, Middlesex, Great Britain). Immunotherapy with a humanized antibody may result in increased long-term effectiveness for the treatment of chronic disease situations or situations requiring repeated antibody treatments.

25 Pharmaceutical compositions may be produced as neutral or salt forms. Salts can be formed with many acids, including, but not limited to, hydrochloric, sulfuric, acetic, lactic, tartaric, malic and succinic acids. Compositions can take the form of solutions, suspensions, suppositories, tablets, pills, capsules, sustained release compounds, or powders. Such 30 formulations can contain 10%-95% (w/w) of the active ingredient, preferably

25%-70% (w/w). Pharmaceutical preparations and compositions can also contain one or more physiologically acceptable carrier(s), excipient(s), diluent(s), disintegrant(s), lubricant(s), plasticizer(s), filler(s), colorant(s), dosage vehicle(s), absorption enhancer(s), stabilizer(s), or bactericide(s).

5 The production and formulation of such compositions and preparations are carried out by methods known and practiced in the art.

Following the preparation of pharmaceutical compositions, they may be placed in appropriate containers and labeled for the treatment of indicated conditions. Such labeling can include amount, frequency, and
10 method of administration. Preparations may be administered systemically by oral or parenteral routes. Non-limiting parenteral routes of administration include subcutaneous, intramuscular, intraperitoneal, intravenous, transdermal, inhalation, intranasal, intra-arterial, intrathecal, enteral, sublingual, or rectal.

15 A therapeutically effective amount of a pharmaceutical composition containing one or more AM-binding proteins or peptides is an amount sufficient to reduce, ameliorate, or eliminate a disease or disorder related to abnormal AM levels. An effective amount can be introduced in one administration or over repeated administrations to an individual being
20 treated. Therapeutic administration can be followed by prophylactic administration, after treatment of the disease. A prophylactically effective amount is an amount effective to prevent disease and will depend upon the specific illness and subject. The therapeutically effective dose may be estimated initially, for example, either in cell culture assays or in animal
25 models, usually mice, rats, rabbits, dogs, sheep, goats, pigs, or non-human primates. The animal model may also be used to determine the maximum tolerated dose and appropriate route of administration. Such information can then be used to determine useful doses and routes for administration in humans.

As an alternative approach, host cells can be genetically engineered to carry a nucleic acid encoding an AM-binding protein, or peptide fragment derived therefrom, and then introduced into a subject in need of modulation or reduction of AM levels. Following expression and production of the AM-binding protein by the cell, the so-produced AM-binding protein or peptide can act to bind AM and effect a disease state or disorder associated with abnormal levels of AM. Host cells may be genetically engineered by a variety of molecular techniques and methods known to those having skill in the art, for example, transfection, infection, or transduction. Transduction as used herein commonly refers to cells that have been genetically engineered to contain a foreign or heterologous gene via the introduction of a viral or non-viral vector into the cells. Transfection more commonly refers to cells that have been genetically engineered to contain a foreign gene harbored in a plasmid, or non-viral vector. Host cells can be transfected or transduced by different vectors and thus can serve as gene delivery vehicles to transfer the expressed products into muscle.

Although viral vectors are preferred for gene transfer therapies, cells can be genetically engineered to contain nucleic acid sequences encoding the desired gene product(s) by various methods known in the art. For example, cells can be genetically engineered by fusion, transfection, lipofection mediated by the use of liposomes, electroporation, precipitation with DEAE-Dextran or calcium phosphate, particle bombardment (biolistics) with nucleic acid-coated particles (e.g., gold particles), microinjection, or genetically engineered microorganisms (K. 20 Yazawa et al, 2000, *Cancer Gene Ther.* 7:269-274). Vectors for introducing heterologous (i.e., foreign) nucleic acid (DNA or RNA) into muscle cells for the expression of active bioactive products are well known in the art. Such vectors possess a promoter sequence, preferably a promoter that is cell-specific and placed upstream of the sequence to be expressed. The vectors 25 may also contain, optionally, one or more expressible marker genes for

expression as an indication of successful transfection and expression of the nucleic acid sequences contained in the vector. In addition, vectors can be optimized to minimize undesired immunogenicity and maximize long-term expression of the desired gene product(s) (see Nabel, 1999, *Proc. Natl. Acad. Sci. USA* **96**:324-326). Moreover, vectors can be chosen based on cell-type that is targeted for treatment. For example, vectors for the treatment of tumor or cancer cells have been described (P.L. Hallenbeck et al., 1999, *Hum. Gene Ther.* **10**:1721-1733; T. Shibata et al., 2000, *Gene Ther.* **7**:493-498; M. Puhlmann et al., 2000, *Cancer Gene Ther.* **7**:66-73; N. Krauzewicz et al., 2000, *Adv. Exp. Med. Biol.* **465**:73-82).

Illustrative examples of vehicles or vector constructs for transfection or infection of the host cells include replication-defective viral vectors, DNA virus or RNA virus (retrovirus) vectors, such as adenovirus, herpes simplex virus and adeno-associated viral vectors. Adeno-associated virus vectors are single stranded and allow the efficient delivery of multiple copies of nucleic acid to the cell's nucleus. Preferred are adenovirus vectors. The vectors will normally be substantially free of any prokaryotic DNA and may comprise a number of different functional nucleic acid sequences. An example of such functional sequences may be a DNA region comprising transcriptional and translational initiation and termination regulatory sequences, including promoters (e.g., strong promoters, inducible promoters, and the like) and enhancers which are active in the host cells. Also included as part of the functional sequences is an open reading frame (polynucleotide sequence) encoding a protein of interest. Flanking sequences may also be included for site-directed integration. In some situations, the 5'-flanking sequence will allow homologous recombination, thus changing the nature of the transcriptional initiation region, so as to provide for inducible or noninducible transcription to increase or decrease the level of transcription, as an example.

In general, the encoded and expressed AM-binding protein may be intracellular, i.e., retained in the cytoplasm, nucleus, or an organelle of a cell, or may be secreted by the cell. For secretion, the natural signal sequence present in the AM-binding protein may be retained. Alternately,

5 an AM-binding protein or a fragment derived therefrom can be fused to a signal sequence to allow secretion of the fusion protein.

As previously mentioned, a marker may be present for selection of cells containing the vector construct. The marker may be an inducible or non-inducible gene and will generally allow for positive selection
10 under induction, or without induction, respectively. Examples of marker genes include neomycin, dihydrofolate reductase, glutamine synthetase, and the like. The vector employed will generally also include an origin of replication and other genes that are necessary for replication in the host cells, as routinely employed by those having skill in the art. As an example,
15 the replication system comprising the origin of replication and any proteins associated with replication encoded by a particular virus may be included as part of the construct. The replication system must be selected so that the genes encoding products necessary for replication do not ultimately transform the cells. Such replication systems are represented by
20 replication-defective adenovirus (see G. Acsadi et al., 1994, *Hum. Mol. Genet.* 3:579-584) and by Epstein-Barr virus. Examples of replication defective vectors, particularly, retroviral vectors that are replication defective, are BAG, (see Price et al., 1987, *Proc. Natl. Acad. Sci. USA*, 84:156; Sanes et al., 1986, *EMBO J.*, 5:3133). It will be understood that the
25 final construct may contain one or more genes of interest, for example, one or more genes encoding bioactive AM-binding proteins. In addition, cDNA, synthetically produced DNA, PCR amplified, or chromosomal DNA may be employed utilizing methods and protocols known and practiced by those having skill in the art.

According to one approach for gene therapy, a vector encoding an AM-binding protein is directly injected into the recipient cells (*in vivo* gene therapy). Alternatively, cells from the intended recipients are explanted, genetically modified to encode an AM-binding protein, and 5 reimplanted into the donor (*ex vivo* gene therapy). An *ex vivo* approach provides the advantage of efficient viral gene transfer, which is superior to *in vivo* gene transfer approaches. In accordance with *ex vivo* gene therapy, the host cells are first infected with engineered viral vectors directing the expression of at least one AM-binding protein, suspended in a 10 physiologically acceptable carrier or excipient such as saline or phosphate buffered saline, and the like, and then administered to the host. The desired AM-binding protein is expressed by the injected cells, which thus introduce the AM-binding protein into the host. The introduced AM-binding protein can thereby be utilized to treat or ameliorate a disorder that is related to altered 15 circulating levels of AM.

EXAMPLES

The examples as described herein are intended to further illustrate the present invention and are not intended to limit the invention.

20 **EXAMPLE 1**

Non-radioactive labeling of AM was accomplished by conjugating synthetic AM comprising amino acids 1-52 (Phoenix Pharmaceuticals, Inc., Mountainview, CA) with succinimidyl esters linked to biotin (Pierce Chemical Co., Rockford, IL), fluorescein (Molecular Probes, Inc., Eugene, OR), or dinitrophenol (DNP; Molecular Probes, Inc.). Briefly, 25 100 µg of AM (16 µM) was dissolved in 1 ml of 50 mM sodium bicarbonate, pH 8.5, and the succinimidyl ester was added to yield a final molar concentration of 10:1 (linker:AM). The mixture was incubated with slow agitation for 1 hr at room temperature and the reaction was terminated by 30 the addition of 0.1 M ethanolamine followed by another incubation for 1 hr.

Unincorporated ligand was removed by extracting the AM using reverse phase Sep-Pak C-18 cartridges (Waters, Milford, MA) and eluting the sample with acidic-isopropanol. The extract was lyophilized and reconstituted in 1 ml of TBS (0.05 M Tris-HCl, 0.15 M NaCl), 0.1% alkali-treated casein (ATC), 0.1% Tween 20, and 0.05% Triton X-100, pH 7.4.

5 Labeled AM was stored at 4°C for as long as 3 months without significant loss of activity.

EXAMPLE 2

AM-binding proteins were identified using an *in vitro* screening method. Total proteins derived from human plasma (2 µl) were electrophoretically fractionated on 3-8% Tris-acetate gel (Novex, San Diego, CA) under non-reducing conditions. For the ligand blotting experiment, the proteins were transferred to a 0.2 mm nitrocellulose membrane that was incubated for 15 min with 1% Nonidet P-40 (NP-40). Non-specific binding was further blocked with a 2 hr incubation in TBS containing 0.1% ATC. Incubations with 70 nM AM labeled with biotin, fluorescein, or DNP were carried out overnight at 4°C in blocking buffer containing 0.1% Tween 20. For biotin detection, the ligand blot was incubated with an avidin-biotin-peroxidase complex and the AEC reagent (Vector Laboratories, Inc., Burlingame, CA). For fluorescein and DNP, the ligand blots were incubated 1 hr at room temperature with rabbit anti-fluorescein or anti-DNP IgG, respectively, (1:1000 in incubation buffer; Molecular Probes, Inc.). Anti-rabbit antibody labeled with alkaline phosphatase (1:2000; Dako Corporation, Carpenteira, CA) was added for 0.5 hr, and the blot was developed using 4-nitro blue tetrazolium chloride/5-bromo-4-chloro-3-indolyl phosphate (NBT/BCIP) as the color-substrate solution (Roche Molecular Biochemicals, Indianapolis, IN). Ligand blotting was used to detect AMBP-1, a 120 kD protein subsequently identified as human complement factor H.

20
25

FIG. 1.

EXAMPLE 3

Competition of full-length AM binding to AM-binding proteins was tested by pre-incubating a membrane having plasma derived proteins bound thereto with 7 μ M of unlabeled peptides; e.g., CGRP, amylin, insulin, IGF-1, PAMP, or AM truncated polypeptides (AM₁₋₁₂, AM₁₆₋₂₁, AM₂₂₋₅₂, AM₁₃₋₅₂, AM₃₄₋₅₂) for 6 hr at 4°C. Following this preincubation, labeled full-length AM was introduced and incubated overnight at 4°C. Competition studies indicated that only full-length AM was able to dissociate the AM/AMBP-1 (AM/fH) complex. FIGS. 2 and 3.

EXAMPLE 4

According to the present invention, a plasma-derived polypeptide that showed specific binding to full-length AM was determined to be factor H. Prior to the formal identification of this AM-binding protein as factor H, it was termed "AM-binding protein" or "AMBP-1" by the present inventors. Isolated and purified AMBP-1 was found to have a molecular weight of approximately 120 kD when subjected to SDS-PAGE under nonreducing conditions.

AMBP-1 was purified to homogeneity using HPLC (FIG. 4A) followed by SDS-PAGE under non-reducing conditions (FIGS. 4B and 4C). Preparative reverse phase high performance liquid chromatography (HPLC) was performed using a Delta Pak C18-300 Å column (30 mm x 30 cm, Waters, Tokyo, Japan) and the "System Gold" modular system (Beckman Instruments Inc., Fullerton, CA). Two and one-half milliliters of human plasma were mixed with an equal volume of 10% acetonitrile with 0.2% trifluoroacetic acid, processed through a 0.2 μ m filter and loaded onto the column. After 5 min with 0.1% trifluoroacetic acid in 5% acetonitrile, the column was eluted with a linear gradient of acetonitrile containing 0.075% trifluoroacetic acid from 5% to 60% at a flow rate of 12 ml/min over 60 min. Each fraction (12 ml) was collected, freeze-dried and dissolved in 0.3 ml of TBS 0.1% Tween 20. Fractions were separated by SDS-PAGE and then

stained with Coomassie Blue (FIG. 4B) or tested for the presence of AMBP-1 using the non-radioactive ligand blotting technique (FIG. 4C).

AMBP-1 was identified in fractions number 48-51, and was found to be concentrated in fractions number 48-49 (FIGS. 4B and 4C).

5 Fraction number 49 was electrophoresed by preparative SDS-PAGE and electroblotted to PVDF. The AMBP-1 band was identified by Coomassie brilliant blue R™ staining, excised, and used for amino acid analysis and amino acid sequence analysis.

EXAMPLE 5

10 Amino acid analysis was performed by The Protein/DNA Technology Center at the Rockefeller University, New York. HPLC (NovaPak C18 30 cm-column) with the Waters PicoTag Workstation and a two pump gradient system (model 510) equipped with a model 490 UV multiwavelength detector were used as previously described (F. 15 Gharahdaghi et al., 1992, in *Techniques in Protein Chemistry III* pp. Academic Press, Inc., San Diego, CA, pp 249-260). The results of the AMBP-1 amino acid analysis are shown below in Table 1 in which % AMBP-1 and % fH are compared.

TABLE 1: Amino acid composition of AMBP-1 and factor H

% of total amino acid composition^a

	AMBP-1	fH ^c
Asx ^b	12.7	9.4
Glx ^b	13.2	12.2
Ser	8.1	8.2
Gly	8.3	8.5
His	2.6	2.5
Arg	4.4	4.9
Thr	6.4	6.6
Ala	3.4	3.3
Pro	7.8	8.0

	43	
Tyr	5.8	6.1
Val	6.4	5.5
Met	0.9	1.7
Ile	5.8	5.9
Leu	5.0	5.0
Phe	2.8	2.7
Lys	6.4	7.0

^aSince tryptophan and cysteine are destroyed by the hydrolysis of the protein in 6 M HCl, the calculated composition of factor H corresponds to the 18 amino acids recovered.

^bThe combined values for asparagine and aspartic acid, and glutamine and glutamic acid are expressed as Asx and Glx, respectively.

^cThe amino acid composition of factor H was obtained from the protein sequence database SWISS-PROT (accession number: P08603).

Although the percentage of methionine determined for AMBP-1 was about one-half of the percentage of methionine predicted for factor H, the recovery of methionine from the control protein (bovine serum albumin) 5 was also about half of the expected value. The amino acid composition profile was used to search a sequence database. This search indicated that the amino acid composition profile shared the highest degree of similarity to human complement factor H.

The N-terminal sequence analysis was performed by the 10 Biotechnology Resource Laboratory, Protein Sequencing and Peptide Synthesis Facility, Medical University of South Carolina, Charleston, SC. The sample was subjected to automated Edman degradation using a PE Biosystems Procise 494 Protein Sequencer and a PE Biosystems cLC Microblotter 173, using standard cycles and reagents (P. Edman and G. 15 Begg, 1967, *Eur J Biochem.* 1:80-91; M.W. Hunkapiller et al., 1983 *Methods Enzymol.* 91:399-413).

The Edman degradation results for AMBP-1 were analyzed using the ExPASy Molecular Biology Server protein database. This analysis indicated that the AMBP-1 amino acid sequence showed an 80% homology match with human complement factor H. Secondary amino acids were 5 attributed to autolytic cleavage of the polypeptide. The sequence of the 15 amino acid N-terminus of AMBP-1 was identical to the N-terminus of factor H with the exception of the threonine in position 12.

For Mass Spectrometry analysis, AMBP-1 was run on a SDS-PAGE gel under reducing conditions (5% β -mercaptoethanol). The gel was 10 stained with Coomassie Blue and the AMBP-1 band was excised. In-gel protein digestion and peptide extraction were performed as previously described (J. Rosenfeld et al., 1992, *Anal. Biochem.* **203**:173-179). One-tenth of the extracted protein digest was analyzed by MALDI-TOF on a PerSeptive Voyager-DE STR (PE Biosystems, Foster City, CA) prior to 15 LC/MS. The instrument was operated in reflector mode with the accelerating voltage set to 20000, the laser energy to 2350, the guide wire voltage to 0.05%, and the grid voltage to 95%. The mirror ratio was set to 1:110.

The remainder of the extracted protein digest was injected 20 onto a 0.3 x 100 mm, 5 μ m BetaBasic C18 column (Keystone Scientific, Bellafonte, PA) which had been equilibrated with 10% buffer B in buffer A (buffer A: water with 0.1% formic acid; buffer B: acetonitrile with 10% 1-propanol and 0.1% formic acid). Peptide elution was carried out using a linear gradient progressing from 10% to 60% buffer B over 60 min 25 (Shimadzu Sci. LC10AD/VP pumps and LC10A controller). The eluting peptides were detected by a Finnigan LCQ Mass Spectrometer (ThermoQuest, Finnigan MAT Division, Piscataway, NJ). Peptide sequence data was obtained from the eluting peptides by MS/MS on those ions exceeding a preset threshold of 5×10^4 ions. The operating parameters 30 were as follows: sheath gas flow set to 32, auxiliary gas flow set to 1, spray

voltage set to 4.5 kV, capillary temperature set to 200°C, capillary voltage set to 8.0 volts, and tube lens offset set to -20 volts.

EXAMPLE 6

The isolated AM-binding plasma protein termed AMBP-1 was
5 compared with factor H, based on apparent molecular weight, glycosylation,
AM-binding, and recognition by anti-factor H antibodies.

Human factor H had been characterized as a glycoprotein
having a calculated molecular weight of approximately 150 kD when run
under reducing conditions (in the presence of 5% 2-mercaptoethanol) (R.B.
10 Sim et al., 1982, *Biochem. J.* **205**: 285-93; P.F. Zipfel et al., 1999, *Mol.
Immunol.* **36**:241-8). It was demonstrated that AMBP-1 was glycosylated
using the GelCode® Glycoprotein Staining Kit (Pierce Chemical Co.), and
that AMBP-1 migrated at a molecular weight of approximately 150 kD under
reducing conditions. FIG. 5A.

15 Additionally, ¹²⁵I-labeled AM (Phoenix Pharmaceuticals, Inc.)
exhibited high-affinity binding to immobilized commercially produced factor H
(10⁻⁸ M to 10⁻⁹ M) (FIG. 5C), and Western blot analysis indicated that
commercially produced antibodies to factor H (Quidel Corporation, San
Diego, CA) recognized plasma-purified AMBP-1 (FIG. 5D). For Western blot
20 analysis, proteins were electrophoretically separated on a 3-8% Tris-acetate
(for factor H) or a 4-12% Bis-Tris gel (for AM) under non-reducing
conditions, transferred to a 0.2 µm nitrocellulose membrane, and blocked
with 5% non-fat dry milk in PBS. Following the blocking step, the membrane
was incubated with 1:2000 anti-factor H rabbit antibody (Quidel, San Diego,
25 CA) or 1:4000 anti-AM rabbit antibody (Martinez et al., 1996, *supra*), and
developed using the ECL+Plus Western Blot Detection System (Amersham
Pharmacia Biotech, Piscataway, NJ).

EXAMPLE 7

To analyze the AM/fH interaction, the purified AMBP-1 (factor
30 H) fraction (#48) was analyzed using the non-radioactive ligand-blotting

assay (Example 1). After incubation with fluorescein-labeled AM, the membrane was incubated under different conditions, including PBS; PBS pH 11.5; PBS pH 2.5; 4M NaCl; 4M NaCl pH 11.5; NaCl pH 2.5; 1% SDS; 3M Urea; 3M Guanidine-HCl; 3M NaSCN; 50% ethylene glycol pH 11.5; 50 % ethylene glycol; 1% β -mercaptoethanol. The assay was then developed as described (Example 1). FIG. 6A.

Alternatively, the AM/fH interaction was analyzed using a multi-well assay. In accordance with this assay, a 96-well polyvinylchloride (PVC) plate was coated with factor H (5 ng/well, Sigma, St. Louis, MO). The plate 10 was blocked with a solution containing TBS, 0.1% ATC, and 0.1% Tween 20, and then incubated with fluorescein-labeled AM (50 nM) for 2 hr. Prior to the development of the assay, the wells were incubated for various time periods with PBS or 3M NaSCN pH 7.4. Following this, the plate was washed and incubated with anti-fluorescein polyclonal antibody (1:1000, 15 Molecular Probes) and 125 I-Protein A (Amersham Pharmacia Biotech). Radioactivity was determined using a gamma counter. FIG. 6B.

EXAMPLE 8

To determine factor H and AM distribution following C18 extraction, 1 ml of human plasma was processed through a Sep Pak C18 20 cartridge. The bound and unbound fractions were tested for the presence of factor H by Western blot analysis (FIG. 7A). Alternatively, bound and unbound fractions were tested for the presence of AM by immunoprecipitation of AM followed by Western blot analysis (FIG. 7B). Immunoprecipitation of AM was performed as follows: a 3 ml sample was 25 incubated for 1 hr at 4°C with 1 ml of Protein A-agarose (Life Technologies, Rockville, MD) containing 1 μ M final concentration of each of the following protease inhibitors: Pefabloc (Centerchem Inc., Stamford, CT), Bestatin, and Phosphoramidon (Sigma). The sample was then centrifuged and the supernatant was removed to another tube. The supernatant was incubated 30 with 80 μ l of rabbit anti-AM antibodies (Martinez et al., 1996, *supra*) or rabbit

preimmune serum for 1 hr at 4°C. The antibody mixture was incubated with 80 µl of Protein A-agarose for 30 min. Following this, the immunoprecipitate was collected by centrifugation and the pellet was extensively washed with TBS with 0.1% Triton X-100. The pellet was then resuspended in 100 µl of 5 LDS sample buffer (Novex) and boiled before the Western blot analysis.

For the standard RIA protocol, plasma extraction was performed using reverse-phase Sep-Pak C18 cartridges (Waters) as previously reported (L.K. Lewis et al., 1998, *Clin. Chem.* **44**: 571-577; A. Martinez et al., 1999, *Peptides* **20**:1471-1478). Briefly, cartridges were 10 activated with 80% methanol and washed with 0.9% NaCl. Plasma samples were mixed with an equal volume of phosphate buffer saline containing 0.1% ATC and 0.1% Triton X-100, pH 7.4. Samples were applied to the columns and, after washing twice with 0.9% NaCl, AM was eluted with 80% isopropanol containing 125 mM HCl. Extracts were freeze-dried to remove 15 the organic solvent. Concentrations of AM in the extracts were measured by radioimmunoassay as previously described (Martinez et al., 1999, *supra*).

For the NaSCN-modified RIA protocol, 1 ml of plasma was pre-incubated with an equal volume of 6M NaSCN in PBS with 0.1% ATC and 0.1% Triton X-100 pH 7.4 for 10 min at room temperature. Following this, 20 plasma was extracted through the C18 cartridges and AM levels were quantified. The AM levels measured using NaSCN-modified RIA were twofold higher than those obtained with the standard RIA. Mean and standard deviation values of the three donors were 23.0 ± 4.8 pg/ml (standard RIA) 54.3 ± 8.6 pg/ml (NaSCN-modified RIA). Identical results 25 were obtained using a longer NaSCN pre-incubation step (16 hr at 4°C). Using the NaSCN-modified RIA, recovery of unlabeled AM added to human plasma (200 pg) was $93.9 \pm 18.7\%$ (n=3), whereas recovery of ^{125}I -AM was $82.7\% \pm 4.4\%$ (n=6). The parallel trajectory of the competitive binding curves indicated that the increase in AM levels detected following NaSCN 30 treatment was not attributable to an artifact. FIG. 8.

EXAMPLE 9

For the cAMP assay, Rat-2 fibroblasts were grown in RPMI 1640 containing 10% fetal bovine serum (Life Technologies). Cells were seeded into 24-well plates at 2×10^5 cells/well and incubated for 48 hr at 5 37°C in 5% CO₂. Cells were preincubated in TIS medium (RPMI 1640 plus 10 µg/ml transferrin, 10 µg/ml insulin and 50 nM sodium selenite) for 15 min. Following this, cells were treated for 5 min with AM (Bachem, King of 10 Prussia, PA) and/or factor H (Sigma) in 250 µl of TIS medium containing 1% BSA, 1 mg/ml bacitracin and 100 µM isobutylmethylxanthine. The reaction was terminated by adding an equal volume of ice-cold ethanol. Cyclic AMP 15 was measured using the Biotrac cAMP RIA (Amersham Pharmacia Biotech). FIG. 9A.

Antimicrobial activity was measured using *E. coli* cells (ATCC 35218, Gaithersburg, MD) and a radial diffusion assay as previously 15 described (R. I. Lehrer et al., 1991, *J. Immunol. Methods* **137**:167-173). Briefly, bacteria were incorporated into a thin underlay gel that contained 1% agarose, 2 mM HEPES pH 7.2, and 0.3 mg/ml of trypticase soy broth powder. After polymerization of the gel, small wells (10 µl capacity) were carved in the agar. Test substances were added and allowed to diffuse for 3 20 hr at 37°C . A 10 ml overlay gel composed of 1% agarose and 6% trypticase soy broth powder was poured on top of the previous gel and the plates were incubated for 16 hr at 37°C . The diameters of the inhibition halos were measured to the nearest 0.1 mm and, after subtracting the diameter of the well, were expressed on inhibition units (10 units = 1 mm). The minimal 25 inhibitory concentration (MIC) was estimated by performing a linear regression and determining the x-intercepts. FIG. 9B.

For statistical calculations, MIC values were analyzed by the Student's t test, cAMP values were analyzed with a one-way analysis of variance (ANOVA) and the Tukey's test, and P<0.05 was considered 30 significant.

To measure the cofactor activity of factor H, C3b (28 pmols) was incubated with factor I (0.16 pmols) and factor H (0.16 pmols) in the presence or absence of AM and related peptides for 24 hr at 37°C in a final volume of 50 µl of PBS. Samples were fractionated by SDS-PAGE using 4-
5 12% Tris-Bis gels (Novex) under reducing conditions, and stained with Coomassie Blue. C3b and factor I were purchased from Advanced Research Technologies (San Diego, CA). FIGS. 10A and 10B.

EXAMPLE 10

Northern blot analysis was used to determine the expression of factor H (4.3 kb) and FHL-1 (1.8 kb) message in various human tumor cell lines. A ³²P labeled 854 bp PCR product was used as a probe to detect Factor H and FHL-1 mRNA in the same Northern blot. The sequence information for factor H and FHL-1 was obtained from GenBank Accession Nos. Y00716 (factor H) and X07523 (FHL-1). To obtain the PCR product/probe, PCR was performed of using a sense primer corresponding to positions 555-576 of factor H (5'-CAATGGAACCAGATCGGGATTA (SEQ ID NO:1)) and the antisense primer corresponding to positions 1378-1408 of factor H (5'-GACACGGATGCATCTGGGAGTA (SEQ ID NO:2)). Total RNA from human liver was reverse transcribed to cDNA and used as template.
10
15
20
25
30

The northern gel was loaded with 15 µg total RNA per well (carcinoma samples), or 5 µg total RNA per well (normal skin sample). To confirm equivalent amounts of total RNA loaded in each well, the gel was stained with ethidium bromide and illuminated with UV. FIG. 11.

EXAMPLE 11

For Immunohistochemical analysis, 6 male albino Wistar rats were euthanized with CO₂, and perfused with the fixative (4% paraformaldehyde) through a heart canula. The pancreas from each rat was extracted and immersed in the same fixative for an additional period of 5 h. After dehydratation, the tissues were embedded in paraffin and sectioned according to routine procedures.
25
30

Two commercially available polyclonal antibodies against human factor H were used in conjunction with an avidin-biotinylated peroxidase detection kit (Dakopatts, Glostrup, Denmark). The antibodies were goat anti human factor H (Quidel, San Diego, CA) and rabbit anti 5 human factor H (Serotec, Raleigh, NC). To ensure the specificity of the signal, liquid-phase and solid-phase absorption controls were performed with purified human factor H (Sigma, St. Louis, MI). For liquid-phase absorptions, 10 nmols of factor H were added per ml of optimally diluted antibody for 2 hr at room temperature before incubating the tissue sections.

10 For solid-phase absorptions, factor H was linked to Ultralink Biosupport Medium (Pierce, Rockford, IL), the optimally diluted antibody was exposed to it for 2 hr at room temperature, and the immune complexes removed by centrifugation. FIGS. 12A-12C.

Anti-factor H antibodies were then purified by affinity 15 chromatography column containing solid-phased factor H. Briefly, 1 mg of factor H was covalently coupled to 250 mg UltraLink Biosupport Medium (Pierce) following manufacturer's instructions. Rabbit anti-factor H serum (250 µl) was incubated with the resin for 16 hr in 5 ml of PBS at 4°C. Following this, the resin was packed in a column and washed intensively 20 with PBS. The bound antibody was eluted with 0.1 M citric acid, pH 3.3. The eluate was neutralized with 1 M sodium phosphate pH 8.0, and the buffer changed to PBS by ultracentrifugation in Centricon 50 columns (Amicon, Millipore Corporation, Bedford, MA). Solid-phase absorption of this affinity-purified antibody resulted in a successful quenching of the staining.

25 FIG. 12D.

EXAMPLE 12

Triple immunofluorescence labeling and confocal microscopy was performed as follows. Non-specific binding was blocked with donkey 30 normal serum (Jackson Immunoresearch Laboratories, West Grove, PA). Paraffin sections were incubated in a mixture of three primary antibodies,

including: i) Guinea-pig anti bovine insulin antibodies (1:2,000) (Jackson Immunoreasearch Labs.); ii) affinity purified rabbit anti human factor H antibodies (1:200) described above; and iii) one of the following monoclonal antibodies: anti-somatostatin antibodies (1:10,000), anti-glucagon antibodies (1:1,000), or anti-rat pancreatic polypeptide antibodies (1:500) (all obtained from CURE (UCLA, CA)).

Following overnight incubation at 4°C, the sections were incubated for 1 hr with a mixture of secondary antibodies, including Cy5-conjugated donkey anti-Guinea-pig antibodies (Jackson Immunoresearch Laboratories), Bodipy-conjugated anti-mouse antibodies (Molecular Probes, Eugene, OR), and biotinylated goat anti-rabbit (Dakopatts), all at a final concentration of 1:200. The sections were then incubated with lissamine rhodamine streptavidin (1:200) (Jackson Immunoresearch Laboratories) for 1 h. After extensive washing, the slides were mounted in SlowFade solution (Molecular Probes) and observed with a Zeiss Laser Scanning microscope 510, equipped with four lasers. FIGS. 13A-13L.

EXAMPLE 13

For immunoelectron micrographic studies, 4 male albino Wistar rats were euthanized and pancreatic-extraction was performed as previously described. Small fragments of each pancreas were fixed in 1% glutaraldehyde plus 2% paraformaldehyde in 0.1 M cacodylate buffer, pH 7.2 at 4°C for 2 h. The fragments were then dehydrated, and embedded in the epoxy resin, TAAB-812 (TAAB Labs. England). Ultrathin sections (60-80 nm) were placed on nickel grids, etched for 5 min in 3% H₂O₂, and subjected to double immunogold staining (J. López et al., 1999, *Gen. Comp. Endocrinol.* 115:309-322). Briefly, the sections were incubated in a mixture of both primary antibodies (anti-insulin and anti-factor H) at 4°C overnight. On the following day, the sections were incubated with secondary antibodies conjugated to gold particles (E-Y Laboratories Inc., San Mateo, CA). The large gold particles (20 nm in diameter) localized the anti-insulin antibody,

while the small gold particles (10 nm) localized the anti-factor H antibody. The grids were washed extensively and stained with 5% aqueous uranyl acetate for 15 min, and with lead hydroxide for 7 min. The stained sections were analyzed using a JEOL-1010 electron microscope. FIGS. 14A-14C.

5 **EXAMPLE 14**

The expression of complement factor H in the pancreas was confirmed by RT-PCR analysis. Primers were designed using the sequence of the mouse gene (GenBank accession number M12660): sense primer (1877-1896) 5'-TTGGAATTCTCCTGCCATTC-3' (SEQ ID NO:3) and 10 antisense primer (2644-2663) 5'-ACCTTCCATCTTGCACACC-3' (SEQ ID NO:4). Total RNA was prepared from the liver and pancreas of two Balb/c mice, and reverse transcription was performed using the SuperScript Preamplification System (Life Technologies, Gaithersburg, MD). PCR amplified was performed with eLONGase enzyme mixture (Life 15 Technologies) for 35 cycles, with an annealing temperature of 57°C. The PCR products were cloned into pCR-Blunt II-TOPO (Invitrogen, Carlsbad, CA) and sequenced by Lark Technologies (Houston, TX). RT-PCR results were confirmed by Southern blot analysis. FIGS. 15A and 15B.

EXAMPLE 15

20 Pancreatic islets were isolated from 12 rats as previously described (A. Martínez et al., 2000, *Endocrinology* 141:406-411). Briefly, the heart of each rat was perfused with Hank's balanced salt solution (Sigma) to remove blood from pancreatic vessels. A small canula was inserted into the common biliary duct while the coledoco was sealed. 25 Twenty to twenty-five milliliters of 0.4 mg/ml collagenase XI (Sigma) in Hank's solution was pumped through the canula until the pancreas was fully inflated. The pancreas was removed and incubated at 37°C for 20 min.. Following this, 20 ml of ice-cold Hank's solution was added and the pancreas dissociated by vigorous vortexing. After several washes, the islets

were collected and distributed into 24 well plates using a dissecting microscope, with approximately 70 islets per well.

Freshly isolated islets were incubated in RPMI-1640 (Life Technologies) containing 5.6 mM glucose for 45 min at 37°C. The supernatant was isolated for analysis and the islets incubated in RPMI-1640 containing 20.6 mM glucose plus the potential secretagogues (AM and/or factor H) for another 45 min. The medium was isolated for analysis and the islets were dissolved in ice-cold ethanol to measure cAMP production. Insulin and cAMP were measured by radioimmunoassay with commercial kits (Amersham, Arlington Heights, IL), according to the manufacturer's directions. FIGS. 16A and 16B.

Insulin release was expressed as the ratio between the contents in the high glucose medium divided by the amount in the low glucose medium to allow for variations in the number of secreting cells and/or their secretory efficiency. Graphs represent the mean and standard deviation of three wells per treatment. These experiments were performed three times with comparable results. Two-tailed Student's test was performed to determine statistical significance. P values < 0.05 were considered statistically significant.

20 EXAMPLE 16

Experiments were carried out utilizing radiolabeled AM as ligand. Radioligand blotting assays were performed to establish that AM-BPs were present in the plasma of several animal species and to further determine the specificity of binding between AM and the plasma-derived AM-binding protein on a solid membrane such as nitrocellulose (see T.H. Elsasser et al., 1999, *Endocrinology* 140:4908-4911).

Blood sera and plasma were collected and pooled from calves (4-5 months old), pigs (6-7 months old), goats (2 years old), dogs (6-8 years old), mice (4-6 months old), chickens (6-8 weeks old), and human (adult male) and were aliquoted and stored at -20°C until needed. Initial

chromatographic determination of high molecular weight plasma proteins capable of binding recombinant human ^{125}I -AM (Phoenix Pharmaceuticals, Mountain View, CA) was accomplished using a 0.7 x 30 cm column of Sephadex[®] G-50 superfine (Pharmacia LKB, Uppsala, Sweden) equilibrated
5 with 50 mM phosphate, 50 mM disodium EDTA, 135 mM NaCl, 0.1% Tween 20 and 0.125% alkaline-hydrolyzed casein (AHC; 7), pH 7.2. For gel chromatography studies of solution binding of ^{125}I -AM to plasma proteins, bovine plasma (0.1 or 0.3 ml) was mixed with 50,000 cpm of ^{125}I -AM, incubated at 4°C overnight and applied to the column with phosphate
10 buffered saline in a final volume of 0.5 ml. One milliliter fractions were collected using gravity flow and the radioactivity was quantified.

Further detection and characterization of AM-BPs were performed using radioligand blotting procedures similar to those described for IGF binding proteins in plasma (P. Hossenlopp et al., 1986, *Anal. Biochem.* 154:138-146). Briefly, plasma or serum from the ten species was diluted 1:5 with water and further mixed 1:2 with SDS Tris-glycineglycerol loading buffer and incubated for 10 min at 70°C. Plasma or serum volumes equivalent to 1.5 μl were loaded onto 10% acrylamide gels under non-reducing conditions. Proteins were separated at 115 v for 0.5 hr and
20 transferred to 0.2 μm nitrocellulose using a semi-moist transfer (Bio-Rad Hercules, CA) at 18 v for 42 min. Molecular weight markers (29 kD, 43 kD, 66 kD, 78 kD, and 116 kD) were added to complementary lanes on each gel to assess the relative sizes of the resolved proteins.

Nitrocellulose blots were incubated for 15 min with 1.5% NP-40
25 (Sigma, St. Louis, MO). Nonspecific binding was further blocked with a 4 hr incubation of the nitrocellulose in phosphate buffered saline containing either 1% BSA (RIA Grade, SIGMA, St. Louis, MO) or 1% AHC. Incubations with radiolabeled recombinant human ^{125}I -AM were performed overnight at room temperature with agitation, with 80,000 CPM ^{125}I -hAM per 5 ml protein
30 blocking matrix. The next morning, the nitrocellulose was washed 10 min

with 0.2% NP-40, and further washed 4 times for 15 min per wash with phosphate buffered saline. Following air drying, the nitrocellulose was loaded into a phosphoimaging cassette (Molecular Dynamics, Sunnyvale, CA) overnight for analysis of resolved band images. Additional

5 characterization of the banding patterns was accomplished by autoradiography of the blots on Kodak AR-5 film with a 4-day exposure at -80°C. Removal of apparent binding proteins from plasma was accomplished using the C-18 reverse-phase Sep-Pak® RIA preparatory technique (A. Martínez et al., 1997, *Endocrinology* 138:5597-5604). FIG. 17.

10 Competitive displacement of ^{125}I -AM from separated proteins on nitrocellulose ligand blot strips was performed to assess the specificity of the AM binding to the AM-binding protein. Blot strips were incubated in buffer containing 0.1% AHC, ^{125}I -AM, and concentrations of recombinant human AM ranging from 10^{-10} to 10^{-6} M. The intensity of the band images
15 was resolved by phosphoimaging densitometry. Additional aspects of AM specificity for the binding protein(s) were assessed by co-incubating ^{125}I -AM overnight with authentic AM₁₋₅₂ (10^{-6} M) or AM₁₋₁₂, AM₁₃₋₅₂, AM₃₄₋₅₂ peptide fragments, or amylin, CGRP or insulin used at 10^{-5} M. The results showed a linear competitive displacement of ^{125}I -AM from the AMBPs in the presence
20 of increasing concentrations of non-labeled synthetic AM (Elsasser et al., 1999, *supra*).

EXAMPLE 17

A comparison of ^{125}I -AM blot binding patterns in bovine plasmas from healthy and parasitized calves was used to determine whether
25 plasma AM-BP content was affected by the health status of a subject (Elsasser et al., 1999, *supra*). Nitrocellulose transfer blots of plasma proteins from normal healthy 4 month old calves (n=3) and calves infected with a vascular endothelial-resident parasite (*Sarcocystis cruzi*), (n=4), were probed with ^{125}I -AM. Plasma from infected calves was obtained on day 30
30 post oral inoculation (250,000 oocysts) at the peak of expression of clinical

signs of the acute phase response associated with the eruption of schizonts from the endothelium (T.H. Elsasser et al., 1988, *J. Endocrinol.* **116**:191-200).

Data on the effects of parasitic infection on the amount of AM-BP in calf plasma were statistically analyzed using an analysis of variance approach based on the general linear models procedure of SAS (SAS 1986 General Linear Models Procedure: SAS for personal computers, SAS Institute, Cary, NC).

Autoradiograms of proteins transferred to nitrocellulose and probed with radiolabeled ^{125}I -hAM demonstrated that plasma from all of the species tested contained at least one protein that bound to the AM tracer, i.e., an AM-binding protein. This AM-BP was a high molecular weight protein of about 120-130 kD, more specifically, 120 kD. In plasma from the calf, goat, sheep, and to a lesser extent dog, an additional band of 140 kD was observed.

In addition, ^{125}I -AM ligand blotting demonstrated that the content of specific AM-binding proteins was decreased by as much as 68% in plasma of calves experiencing the acute phase response to a parasitic challenge ($P<0.03$). Because both healthy and parasitized calves had similar plasma concentrations of total proteins, albumin and globulins, the measured decrease in AM-BPs in the parasitized calves reflects a specific response in this protein class to the onset of disease. FIG. 18.

EXAMPLE 18

A small molecule combinatorial library can be obtained from the Developmental Therapeutics Program at NIH. This library provides a total of 250,000 compounds that have a molecular weight below than M_r 600. The compounds are organized in 8,000 "families" with similar structural characteristics. One member of each family can be assayed for binding to AM, factor H, or AM/fH. If binding is detected, the remaining family members can be analyzed to identify compounds with increased binding

affinity for AM, factor H, or AM/fH. In the initial screen, 96-well plates can be coated with AM, factor H, or AM/fH (synthetic, recombinant, or isolated), and then incubated with various test compounds. After extensive washing, the presence of AM, factor H, or AM/fH detected with corresponding antibodies, 5 followed by a secondary antibody and ^{125}I -protein A. Following this, individual wells can be scanned with a gamma counter. Wells containing AM-, factor H- or AM/fH-binding compounds can be identified by an increase in signal compared the negative control (no compound added).

Compounds that are identified by their binding activity can be 10 further analyzed using various bioassays. For example, compounds can be tested using the fibroblast cell line, Rat-2, which expresses specific AM receptors (H.A. Coppock et al., 1999, *Biochem. J.* **338**:15-22.). To test AM-binding compounds, the cells can be incubated with AM alone (negative control) or AM plus the compound. To test fH- binding compounds, the cells 15 can be incubated with AM alone, factor H alone, AM plus factor H (negative controls), or AM plus factor H and the compound. To test AM/fH-binding compounds, the cells can be incubated with AM/fH alone (negative control) or AM/fH plus the compound. Following this, the cAMP response of the cells can be analyzed by radioimmunoassay as previously described (A. 20 Martínez et al., 1997, *Endocrinology* **138**:55977-56048; M.J. Miller et al., 1996, *J. Biol. Chem.* **271**:23345-23351; A. Martínez et al., 1999, *Peptides* **20**:1471-1478).

Dose-response curves can be plotted and used to identify 25 compounds that inhibit AM-mediated cAMP production (compared to the negative control(s)). The curves can also be used to identify high-affinity compounds. High-affinity compounds can then be used as models for secondary combinatorial libraries that can include ~ 500 compounds. Such libraries can be assembled from chemically modified forms of the original compound. These new compounds can be tested in the same manner

described above to refine the screening process and identify a potent inhibitor of AM function, factor H function, or AM/fH interaction.

All patent applications, patents, texts, and literature references
5 cited in this specification are hereby incorporated herein by reference in their entirety to more fully describe the state of the art to which the present invention pertains.

As various changes can be made in the above methods and
10 compositions without departing from the scope and spirit of the invention as described, it is intended that all subject matter contained in the above description, shown in the accompanying drawings, or defined in the appended claims be interpreted as illustrative, and not in a limiting sense.

WHAT IS CLAIMED IS:

1. A method for measuring adrenomedullin levels in a sample, comprising:
 - i) incubating the sample with a chaotropic agent to dissociate adrenomedullin and factor H;
 - ii) fractionating the sample to obtain a peptide fraction; and
 - iii) quantifying adrenomedullin levels in the peptide fraction of the sample.
- 10 2. The method according to claim 1, wherein the sample is selected from the group consisting of plasma, serum, cell lysate, and tissue extract samples.
- 15 3. The method according to claim 1, wherein the chaotropic agent is sodium thiocyanate.
4. The method according to claim 1, wherein the sample is fractionated by a C18 matrix.
- 20 5. The method according to claim 1, wherein the adrenomedullin levels are quantified in step (iii) by radioimmunoassay analysis.
6. A method for monitoring adrenomedullin concentrations in a subject having a condition associated with elevated levels of adrenomedullin, wherein the condition selected from the group consisting of heart disease, pulmonary disease, liver cirrhosis, cancer, diabetes, sepsis, and inflammation, comprising:
 - i) incubating a sample from the subject to be analyzed with isolated factor H, or an adrenomedullin-bindable peptide fragment

thereof, under conditions allowing binding of factor H or the factor H peptide to adrenomedullin in the sample

ii) quantifying the amount of adrenomedullin bound to factor H or the factor H peptide; and

5 iii) comparing the levels of adrenomedullin in the subject with levels of adrenomedullin in normal individuals.

7. The method according to claim 6, wherein the monitoring of adrenomedullin concentration assists in diagnosing a condition, determining
10 the severity of a condition, or following the recovery course of a condition.

8. The method according to claim 6, wherein the sample is selected from the group consisting of plasma, serum, cell lysate, and tissue extract samples.
15

9. The method according to claim 6, wherein the cancer is selected from the group consisting of urinary bladder, urethral, renal, rectal, colon, small intestine, gastric, esophageal, salivary gland, gallbladder, liver, breast, vaginal, endometrial, ovarian, cervical, prostate, skin, lung, and brain
20 cancers.

10. The method according to claim 6, wherein the factor H or the factor H peptide comprises a label.
25 11. The method according to claim 10, wherein the label is non-radioactive.

12. The method according to claim 10, wherein the label is an enzyme.
30

13. The method according to claim 6, wherein prior to step (i), the sample is fractionated to obtain a peptide fraction.

14. The method according to claim 13, wherein the sample is fractionated using a C18 matrix.

15. The method according to claim 13 or 14, wherein prior to fractionation, the sample is incubated with a chaotropic agent.

10 16. The method according to claim 15, wherein the chaotropic agent is sodium thiocyanate.

17. A kit for detecting adrenomedullin concentrations in a sample comprising:

15 i) factor H or an adrenomedullin-binding peptide fragment thereof; and

ii) at least one component to facilitate or detect AM-binding.

20 18. The kit according to claim 17, wherein the peptide comprises a label.

19. The kit according to claim 17, wherein the label is non-radioactive.

25 20. The kit according to claim 17, wherein the label comprises an enzyme.

21. A method of identifying an antagonist agent, comprising:

i) contacting adrenomedullin with a test compound under conditions that allow the adrenomedullin to bind to the test compound, and thereby form a stable association;

ii) detecting the adrenomedullin-test compound association of step (i); and

iii) contacting the adrenomedullin-test compound association with factor H under conditions that allow the test compound to inhibit the complexation of adrenomedullin and factor H, wherein inhibition indicates antagonist activity.

10 22. A method of identifying an antagonist agent, comprising:

i) contacting factor H with a test compound under conditions to allow the factor to bind to the test compound, and thereby form a stable association;

15 ii) detecting the factor H-test compound association of step (i);

iii) contacting the factor H-test compound association with adrenomedullin under conditions that allow the test compound to inhibit the complexation of adrenomedullin and factor H, wherein inhibition indicates 20 antagonist activity.

25 23. A pharmaceutical composition comprising an antagonist agent identified according to the method of claim 21 or 22, and a physiologically acceptable carrier, excipient, or diluent.

24. A method of inhibiting the growth or proliferation of cancer or tumor cells, comprising:

i) administering to cancer or tumor cells an antagonist agent identified according to the method of claim 21 or 22 in an amount

effective to inhibit complexation of adrenomedullin and factor H, and a physiologically acceptable carrier, excipient, or diluent; and

ii) inhibiting the growth or proliferation of the cancer or tumor cells.

5

25. The method according to claim 24, wherein the cells are selected from the group consisting of urinary bladder, urethral, renal, rectal, colon, small intestine, gastric, esophageal, salivary gland, gallbladder, liver, breast, vaginal, endometrial, ovarian, cervical, prostate, skin, lung, and brain cancer or tumor cells.

10

26. An isolated and purified protein complex comprising adrenomedullin and factor H.

15 27. An antibody specifically reactive with an epitope formed by the complexation of adrenomedullin and factor H, or a complex-specific fragment thereof.

28. The antibody according to claim 27, wherein the antibody is a
20 monoclonal antibody.

29. A pharmaceutical composition comprising the antibody according to claim 28, and a physiologically acceptable carrier, excipient, or diluent.

25

30. A method of isolating adrenomedullin/factor H protein complexes from a biological sample, comprising:

i) providing a solid support having bound thereto the antibody of claim 27 or 28;

30

ii) contacting said support with the sample; and

iii) eluting the adrenomedullin/factor H complex that binds to the antibody bound to the support.

31. The method according to claim 30, wherein the biological sample is selected from the group consisting of plasma, serum, cell lysate, and tissue extract samples.

32. A method for detecting adrenomedullin/factor H protein complexes in a biological sample, comprising:

10 i) incubating the sample with the antibody of claim 27 or
28 under conditions allowing binding of the antibodies to the adrenomedullin/factor H complex in the sample; and
ii) determining binding of the antibodies to the adrenomedullin/factor H complex.

15 33. The method according to claim 32, wherein the sample is selected from the group consisting of plasma, serum, cell lysate, and tissue extract samples.

20 34. The method according to claim 32, wherein the antibodies are labeled.

35. The method according to claim 32, wherein the antibodies are non-radioactively labeled.

25 36. The method according to claim 32, wherein the antibodies are labeled with an enzyme.

30 37. A kit for detecting adrenomedullin/factor H complex concentrations in a sample comprising:

- i) an antibody specifically reactive with an epitope formed by the complexation of adrenomedullin and factor H, or a complex-specific fragment thereof; and
- ii) at least one component to facilitate or detect
5 adrenomedullin/factor H complex-binding.

38. The kit according to claim 37, wherein the antibody comprises a label.

10 39. The kit according to claim 38, wherein the label is non-radioactive.

40. The kit according to claim 38, wherein the label comprises an enzyme.

15 41. A method of inhibiting the growth or proliferation of cancer or tumor cells, comprising:

- i) administering to cancer or tumor cells an antibody specifically reactive with an epitope formed by the complexation of
20 adrenomedullin and factor H or a complex-specific fragment thereof, in an amount effective to inhibit activity of the adrenomedullin/factor H complex, and a physiologically acceptable carrier, excipient, or diluent; and
- ii) inhibiting the growth or proliferation of the cancer or tumor cells.
—

25 42. The method according to claim 41, wherein the cells are selected from the group consisting of urinary bladder, urethral, renal, rectal, colon, small intestine, gastric, esophageal, salivary gland, gallbladder, liver, breast, vaginal, endometrial, ovarian, cervical, prostate, skin, lung, and brain
30 cancer or tumor cells.

43. The method according to claim 41, wherein the antibody is humanized.

5 44. The method according to claim 41, wherein the antibody is recombinant.

45. A method of treating a condition associated with decreased secretion of insulin by pancreatic cells, comprising:

10 i) administering to the pancreatic cells an antagonist agent identified according to the method of claim 21 or 22 in an amount effective to inhibit complexation of adrenomedullin and factor H, and a physiologically acceptable carrier, excipient, or diluent; and

15 ii) increasing the secretion of insulin by the pancreatic cells.

46. The method according to claim 45, wherein the condition is selected from the group consisting of diabetes and pheochromocytoma.

20 47. The method according to claim 45, wherein the condition is type 2 diabetes.

48. A method of treating a condition associated with decreased secretion of insulin by pancreatic cells, comprising:

25 i) administering to the pancreatic cells an antibody specifically reactive with an epitope formed by the complexation of adrenomedullin and factor H or a complex-specific fragment thereof, in an amount effective to inhibit activity of the adrenomedullin/factor H complex, and a physiologically acceptable carrier, excipient, or diluent; and

ii) increasing the secretion of insulin by the pancreatic cells.

49. The method according to claim 48, wherein the condition is
5 selected from the group consisting of diabetes and pheochromocytoma.

50. The method according to claim 48, wherein the condition is type 2 diabetes.

10 51. A method of treating a condition associated with an autoimmune response to pancreatic cells, comprising:

i) administering factor H to the pancreatic cells in an amount effective to inhibit complement-mediated cell lysis, and a physiologically acceptable carrier, excipient, or diluent; and

15 ii) decreasing the autoimmune response to the pancreatic cells.

52. The method according to claim 51, wherein the condition is type 1 diabetes.

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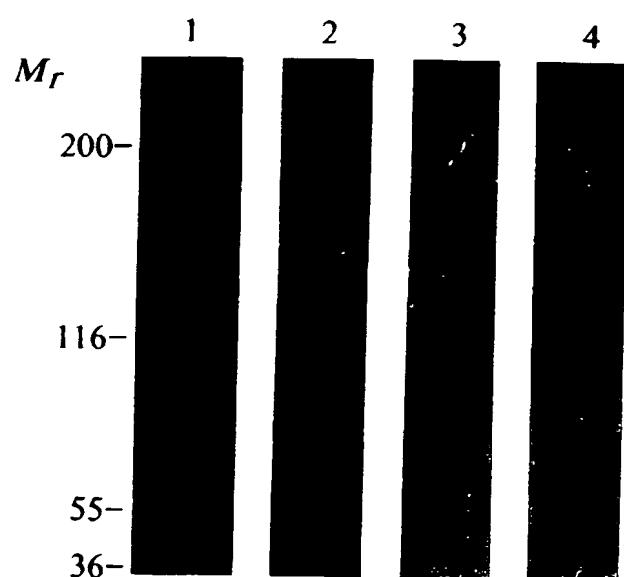


FIG. 1

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f-AM +

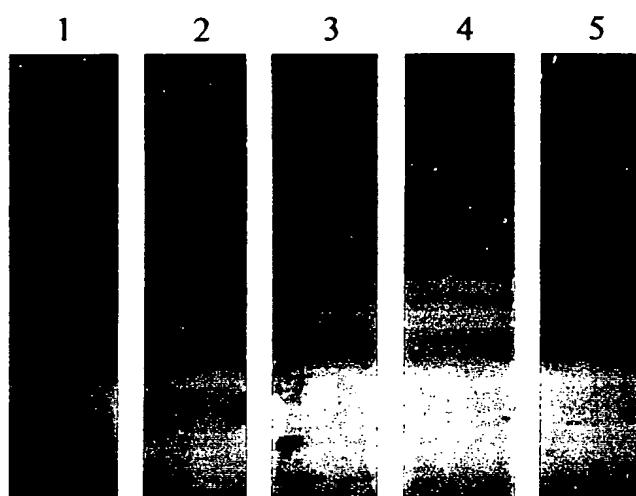


FIG. 2

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f-AM +

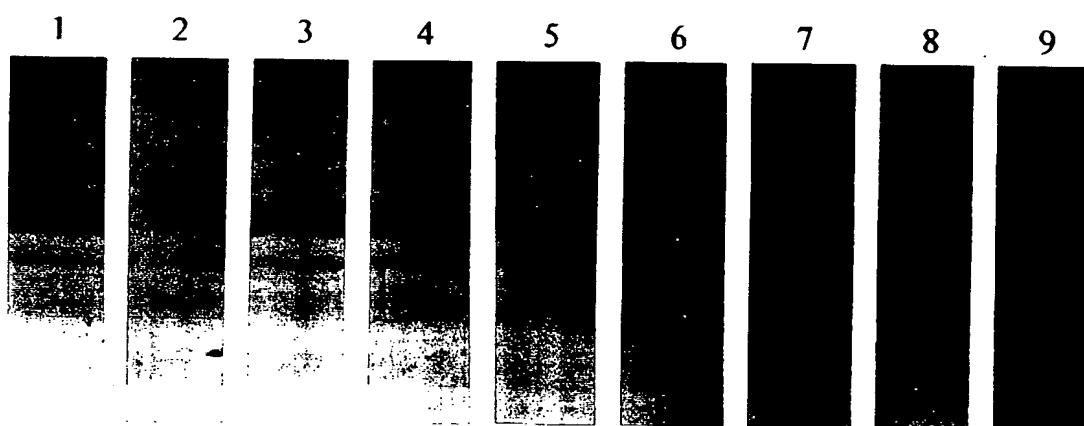
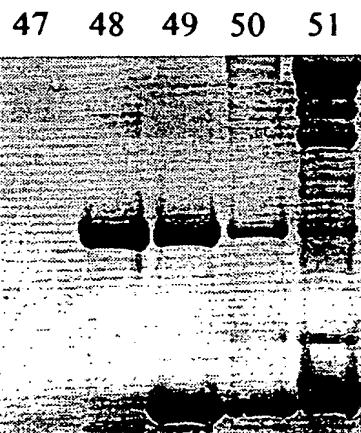
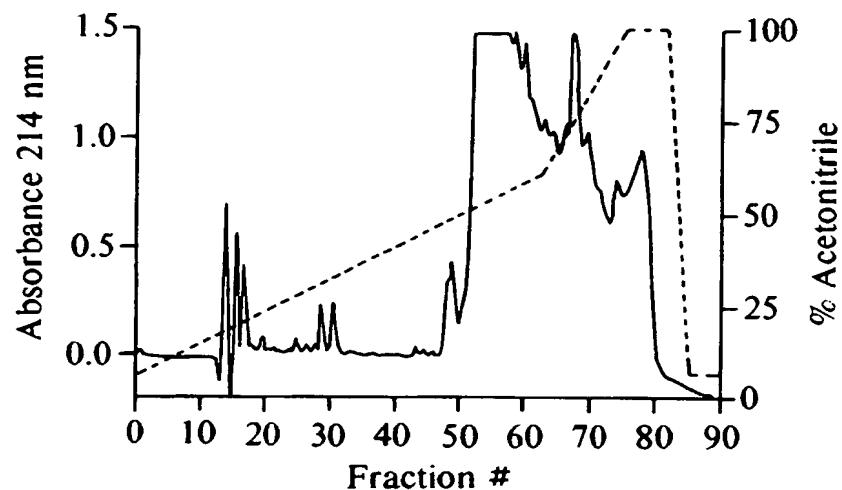
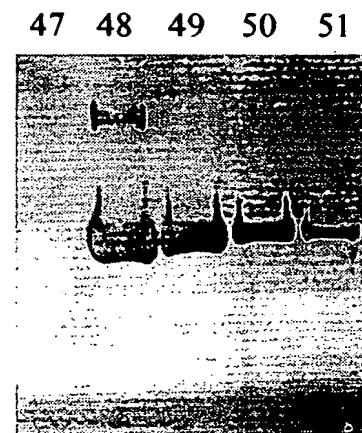


FIG. 3

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FIG. 4A**FIG. 4B****FIG. 4C**

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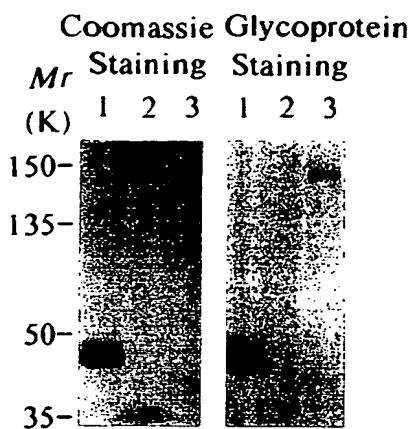


FIG. 5A



FIG. 5B



FIG. 5C

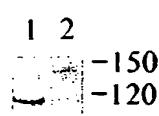
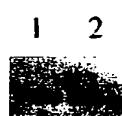


FIG. 5D

FIG. 5E

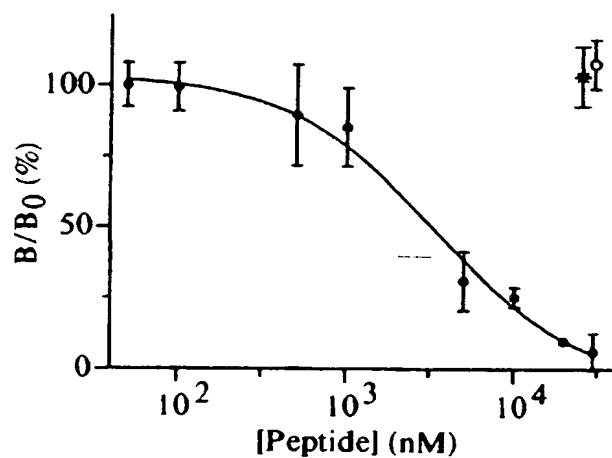


FIG. 5F

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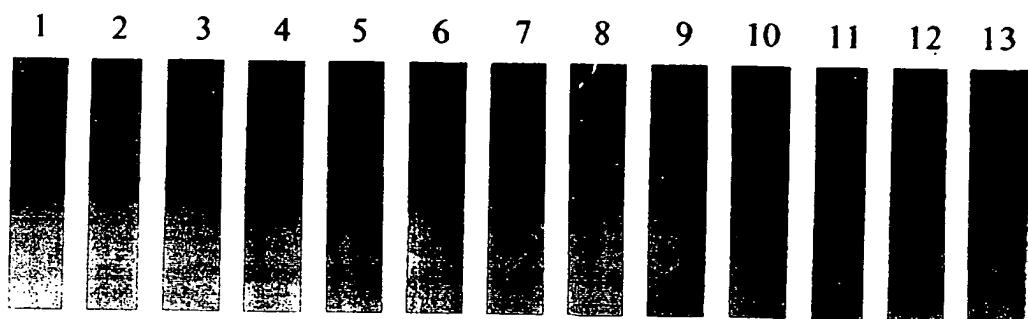


FIG. 6A

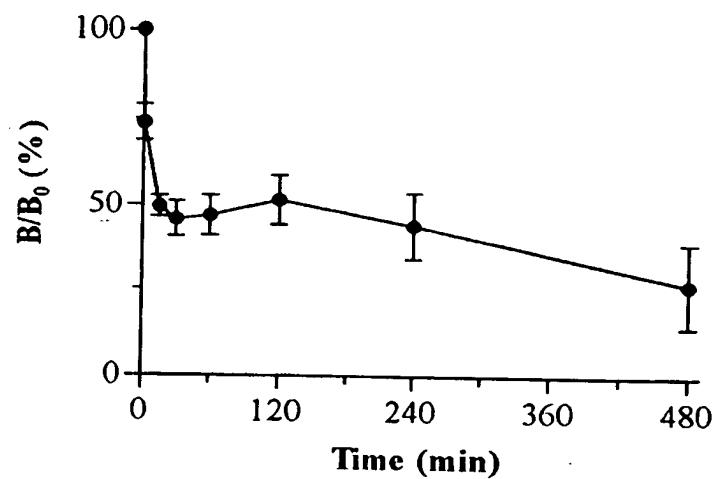
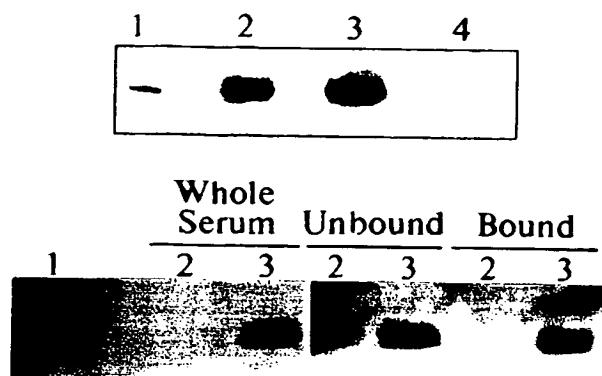
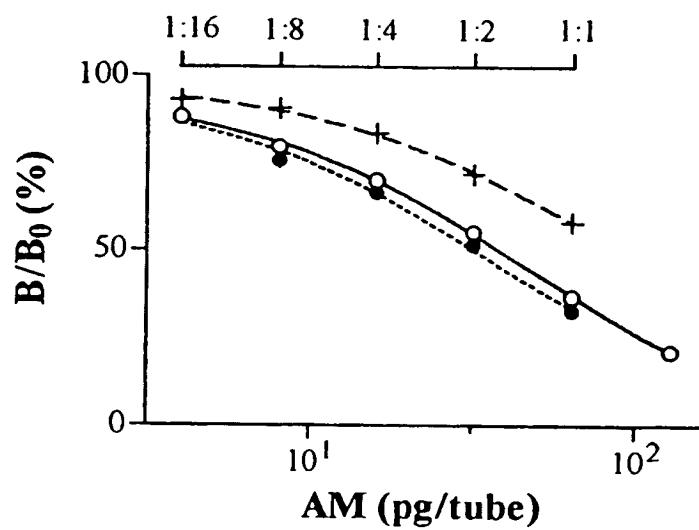


FIG. 6B

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FIG. 7A**FIG. 7B****FIG. 8**

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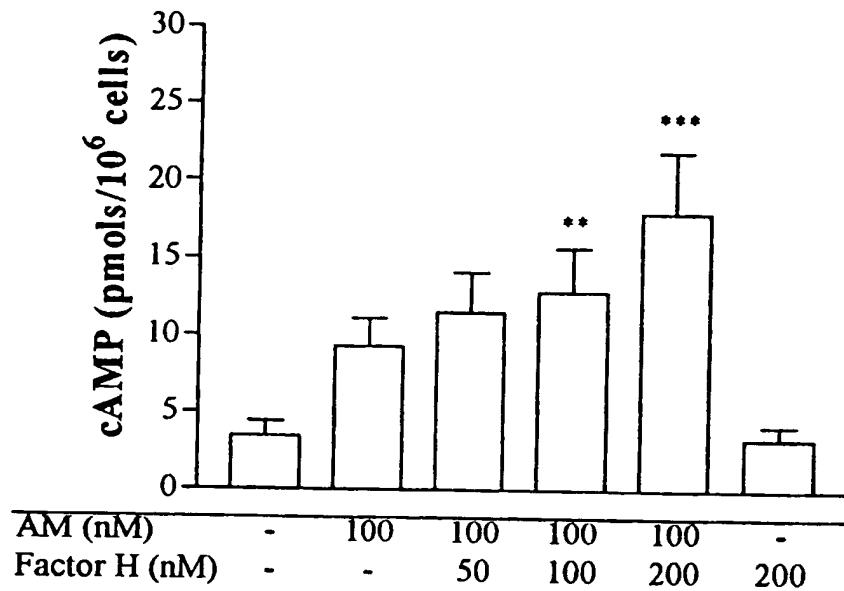


FIG. 9A

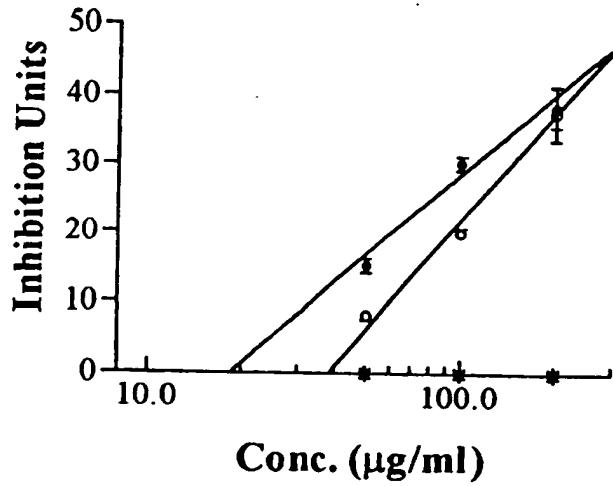


FIG. 9B

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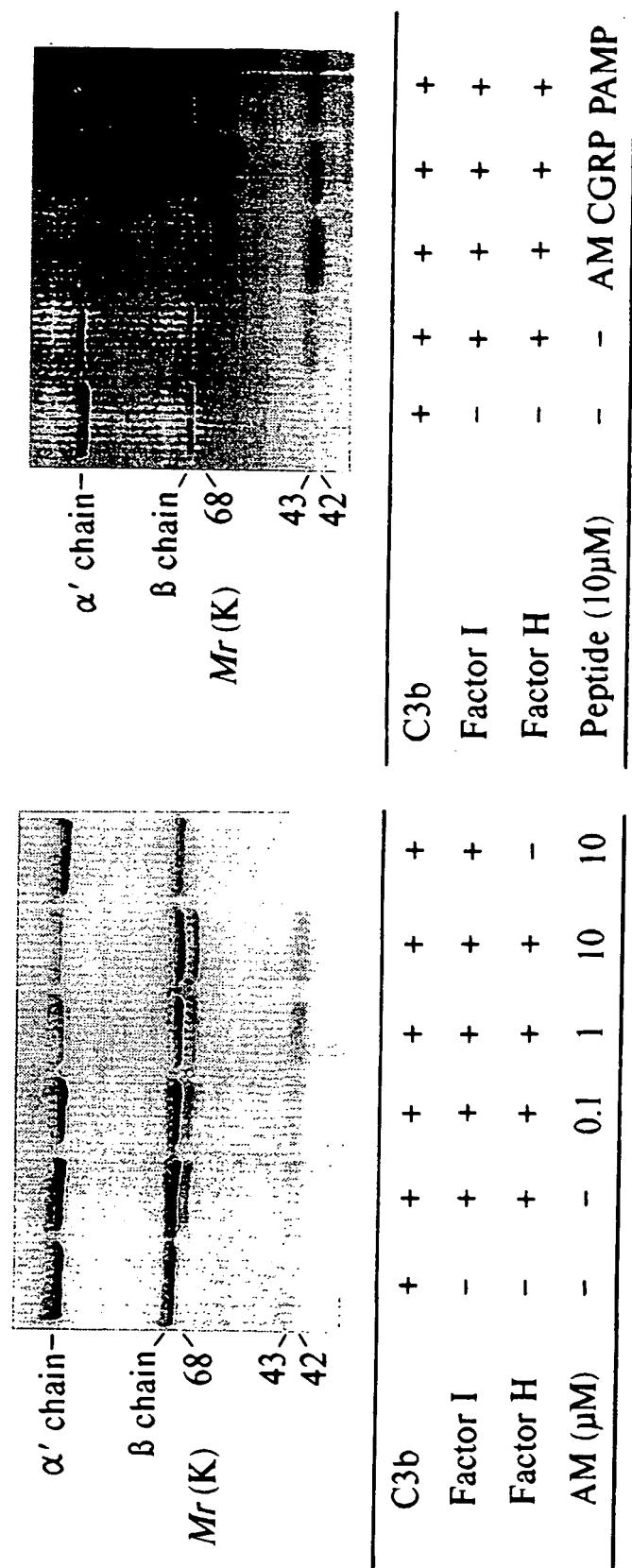


FIG. 10A

FIG. 10B

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FIG. 11A
hCFH Probe

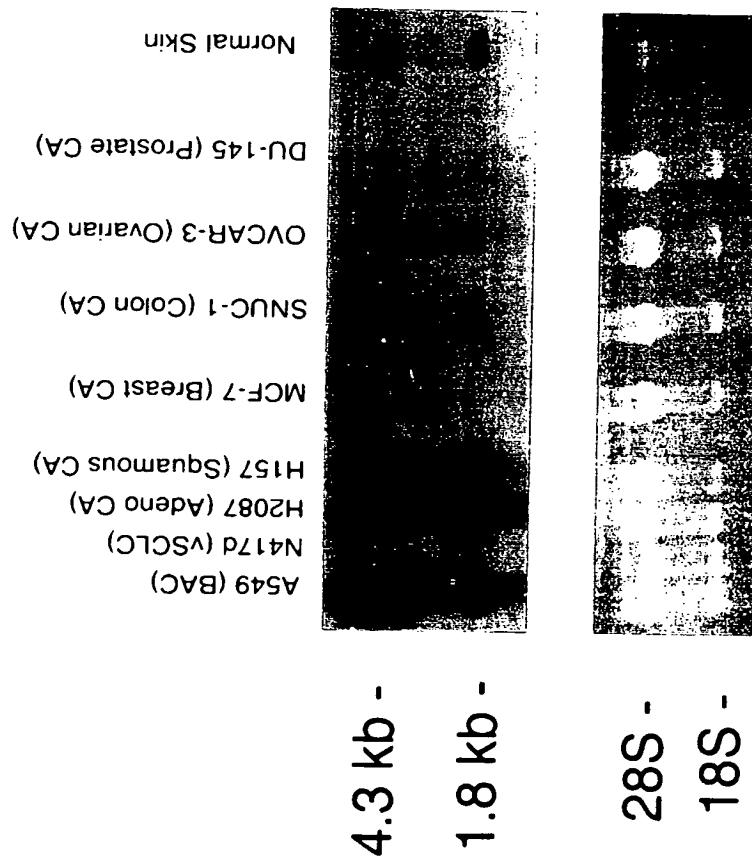


FIG. 11B
UV

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FIG. 12A

FIG. 12B

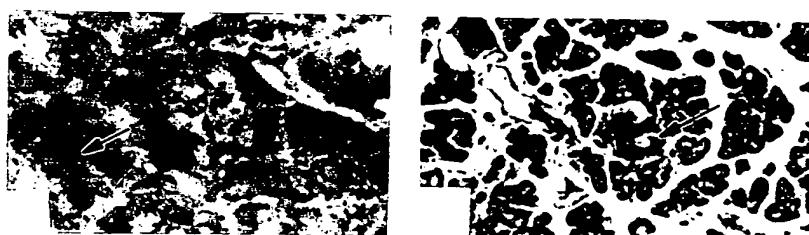


FIG. 12C

FIG. 12D

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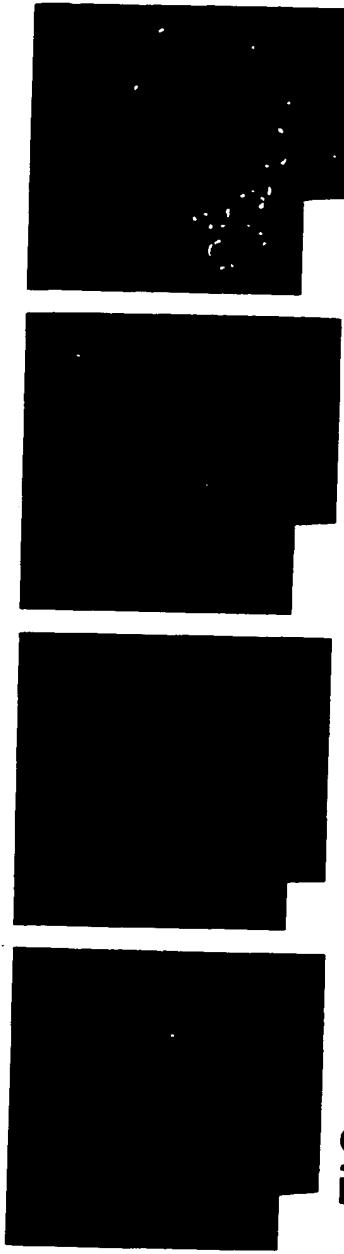


FIG. 13A

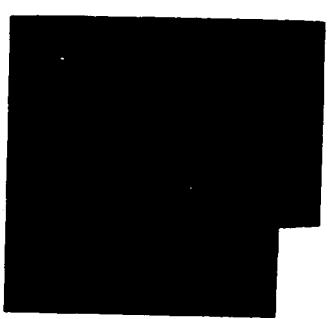


FIG. 13B



FIG. 13C



FIG. 13D

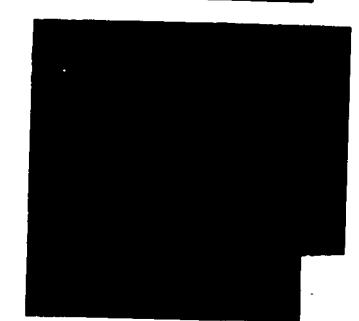


FIG. 13E



FIG. 13F



FIG. 13G



FIG. 13H

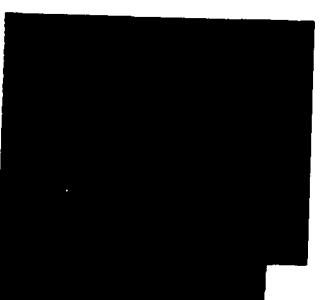


FIG. 13I

FIG. 13L

FIG. 13K

FIG. 13J

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FIG. 14A



FIG. 14B

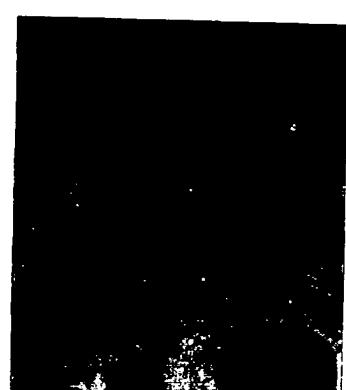


FIG. 14C

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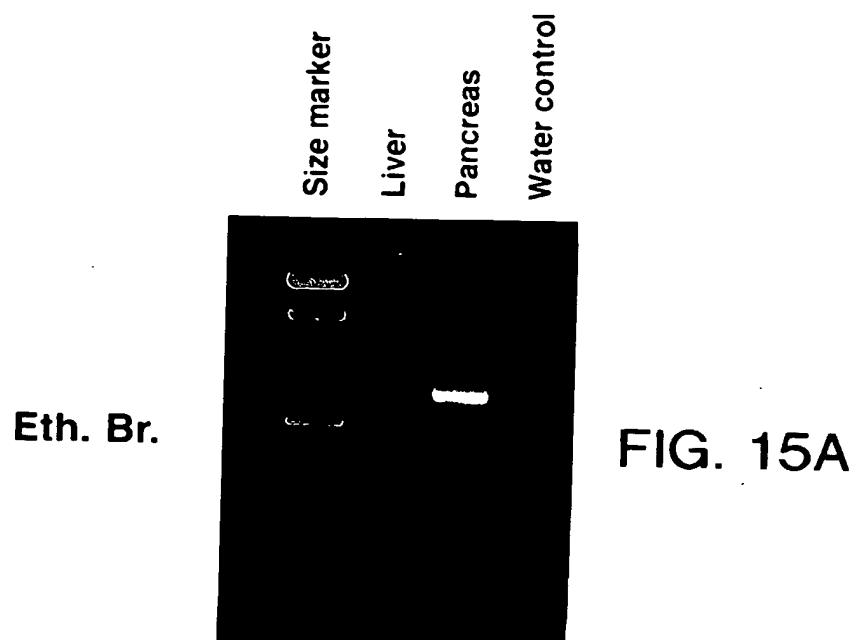


FIG. 15A

Southern

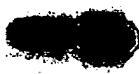


FIG. 15B

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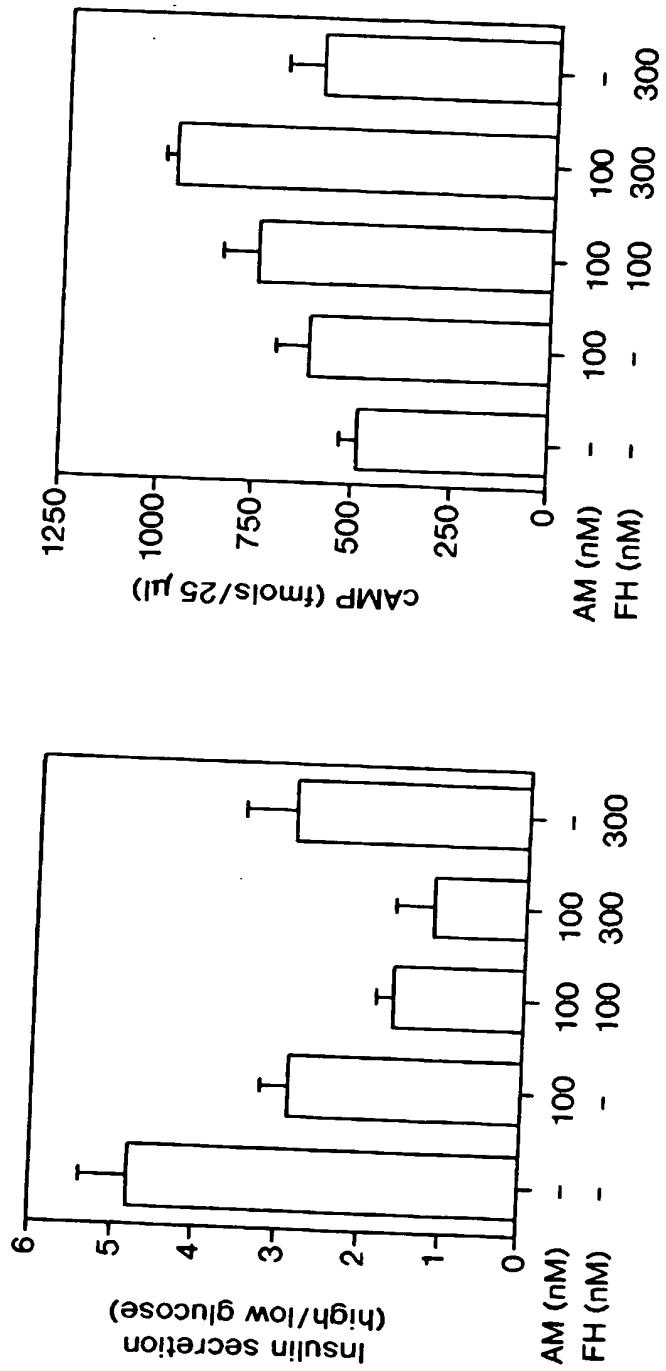


FIG. 16A

FIG. 16B

16/16

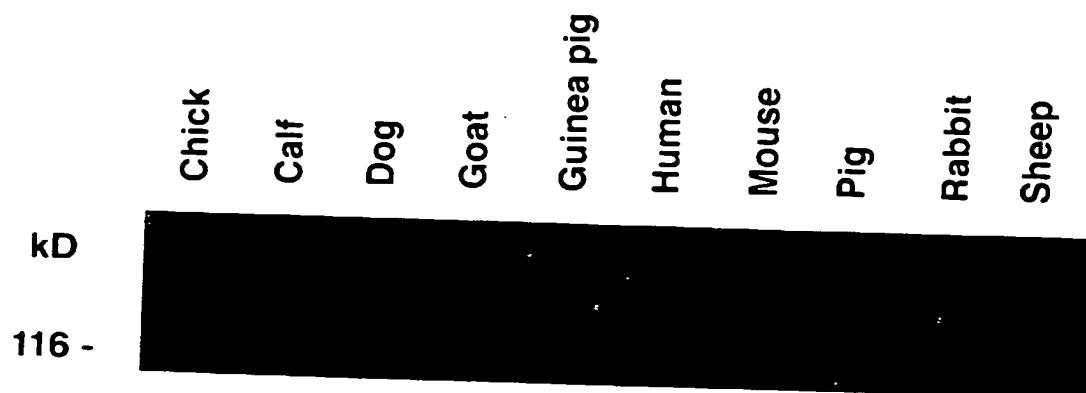


FIG. 17

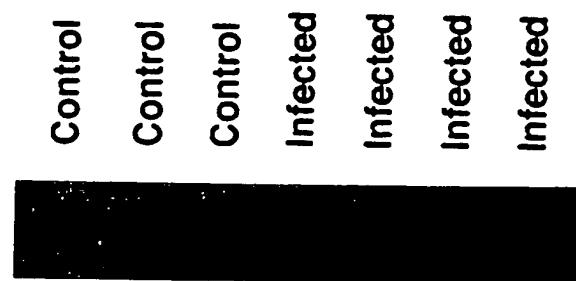


FIG. 18

SEQUENCE LISTING

<110> Cuttitta, Frank
Elsasser, Ted H.
Martinez, Alfredo
Pio, Ruben

<120> DETERMINATION OF AM-BINDING PROTEINS AND THE
ASSOCIATION OF ADRENOMEDULLIN (AM) THEREWITH

<130> 2026-4316PC

<140>
<141>

<150> 60/153,397
<151> 1999-09-10

<160> 4

<170> PatentIn Ver. 2.1

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<211> 22
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<213> Artificial Sequence

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<223> Description of Artificial Sequence: PCR primer

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<223> Description of Artificial Sequence: PCR primer

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<223> Description of Artificial Sequence: PCR primer

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<210> 4

<211> 20

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR primer

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20

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/24722

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 G01N33/74 C07K16/00 A61K38/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHEDMinimum documentation searched (classification system followed by classification symbols)
 IPC 7 G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, BIOSIS, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MARTINEZ ALFREDO ET AL: "Regulation of insulin secretion and blood glucose metabolism by adrenomedullin." ENDOCRINOLOGY, vol. 137, no. 6, 1996, pages 2626-2632, XP000982198 ISSN: 0013-7227 the whole document ---	1-52
A	WO 97 07214 A (UNSWORTH EDWARD J ;MACRI CHARLES (US); US HEALTH (US); CUTTITTA FR) 27 February 1997 (1997-02-27) the whole document ---	1-52
		-/-

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

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- *Z* document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
16 February 2001	26/02/2001
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel: (+31-70) 340-2040, Tx. 31 651 epo nl. Fax: (+31-70) 340-3016	Authorized officer Pellegrini, P

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 00/24722

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MARTINEZ ALFREDO ET AL: "Expression of adrenomedullin and its receptor in normal and malignant human skin: A potential pluripotent role in the integument." ENDOCRINOLOGY, vol. 138, no. 12, December 1997 (1997-12), pages 5597-5604, XP000982199 ISSN: 0013-7227 cited in the application the whole document ---	1-52
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P, A	PIO RUBEN ET AL: "Isolation and characterization of a human plasma adrenomedullin binding protein. Analysis of its expression in human tumor cell lines." PROCEEDINGS OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH ANNUAL, no. 41, March 2000 (2000-03), page 152 XP000982162 91st Annual Meeting of the American Association for Cancer Research.; San Francisco, California, USA; April 01-05, 2000, March, 2000 ISSN: 0197-016X abstract --- -/-	1-52

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 00/24722

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/24722

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